



PATENT DOCKET 709

THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of

Paul J. Carter et al.

Serial No. 07/715272

Filed: June 14, 1991

For: Immunoglobulin Variants

) Group Art Unit: 1806

) Examiner: L. FEISEE

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DECLARATION OF ROBERT F. KELLEY PURSUANT TO 37 CFR §1.132

Honorable Commissioner of Patents
and Trademarks
Washington, D.C. 20231

Sir:

I, ROBERT F. KELLEY, do hereby declare as follows:

1. I received my Ph.D. in Biochemistry in 1984 from the University of Iowa. Following my Ph.D, I was a NIH postdoctoral fellow in the Department of Molecular Biophysics & Biochemistry at Yale University from July 1984 to December 1985. In 1986, I joined the Biocatalysis Department at Genentech, Inc. as an Associate Scientist. In September 1988, I was promoted to Scientist and I am employed in that capacity at present. (The Biocatalysis Department has been renamed "Protein Engineering"). I am the author or co-author of 22 publications relating to the 3-D structures and folding of various proteins. A copy of my curriculum vitae is attached as Exhibit "A".

2. I understand that the Patent Office has rejected the above application on the basis that the application as filed does not provide sufficient disclosure to enable a skilled biochemist to carry out the method of claim 1 because the Examiner believes no clear guidance exists in the specification to allow a skilled biochemist to make the "consensus human variable domain" and substitute an import (i.e. non-human) Complementary Determining Region (CDR) amino acid sequence for the corresponding human CDR amino acid sequence, as set forth in claim 1. I further understand that the Office considers that

the only guidance in the specification with regards to the substitutions is the amino acid sequences of SEQ ID NO: 3 and 4.

3. I have read the above application, the Office Action date May 19, 1992 (Paper # 17) rejecting the claims of the application, and the proposed amendment of the claims in response to the rejection. In my opinion, the skilled biochemist could have readily carried out the method of claim 1 in order to make a humanized antibody, using the general knowledge available in the field on and before June 14, 1991, and the information given in the above application. The bases for my opinion are given in paragraphs 4 to 7 below.

4. Claim 1 relates to a method of making a humanized antibody. Steps a and b of claim 1, as amended, discuss identification of the CDR amino acid sequences of a non-human import antibody (to be humanized) and a consensus human variable domain of a human immunoglobulin subgroup. The consensus human variable domain constitutes an amino acid sequence comprising the most commonly occurring amino acids at each position in the variable domain of a particular human immunoglobulin subgroup as defined by Kabat *et al.*, *Sequences of Proteins of Immunological Interest*, Fourth Edition, U.S. Dept. of Health & Human Services, pubs., (1987), a copy of which is attached as Exhibit "B". The immunoglobulin subgroups referred to in Kabat *et al.* were grouped according to the amino acid sequence homology between human immunoglobulin *variable* domains, and the most commonly occurring amino acids at each position in the variable domain for each subgroup were identified (i.e. the "consensus human variable domain"). The skilled biochemist could have used the consensus human variable domains of the light chain and heavy chain subgroups having the greatest number of sequences (i.e. light chains kappa subgroup I and heavy chains subgroup III) as disclosed in Kabat *et al.* (see page 17, first paragraph of the specification) to humanize the non-human antibody of interest. Alternatively, the skilled biochemist could have chosen the consensus human variable domain of another human immunoglobulin subgroup as defined in Kabat *et al.* (i.e. the consensus human variable domain for human kappa light chains subgroups II to IV, human lambda light chains subgroups I to VI, or human

heavy chains subgroups I or II [see pages 41-76 and 160-167 of Kabat *et al.*]). Therefore, the skilled biochemist could have elected to use a consensus human variable domain other than those defined as SEQ ID NO: 3 & 4 on page 17 of the above application, as the consensus human variable domains for other subgroups were compiled in Kabat *et al.* Page ix of Kabat *et al.* identifies the residues forming the CDR regions of heavy and light chain variable domains tabulated from human and mouse variable domains. Kabat *et al.* have adopted standardized numbering for each of the residue locations. Accordingly, the skilled biochemist could have identified the CDR regions of the consensus human variable domain and the import variable domain using the teachings of Kabat *et al.* Alternatively, the structural definition of Chothia *et al.*, *J. Mol. Biol.*, **196**: 901-917 (1987) (see page 16, third paragraph of the specification) could have been adopted to identify the CDR regions of the consensus and import variable domains. Hence, it would have been straightforward for the skilled biochemist to carry out steps a and b of claim 1 using the information provided in the specification.

5. Step c of claim 1 discloses the step of replacing the corresponding human CDR sequence with the import CDR amino acid sequence. This step could have been carried out routinely by the skilled biochemist by manual tabulation or using a computer program such as the ALIGN program, (Dayhoff *et al.*, *Meth. Enzymol.*, **91**:524-545 [1983]) which was available prior to June 14, 1991. Steps a to c of claim 1 would have resulted in the characterization of a primary amino acid sequence encoding a humanized variable domain with import (non-human) CDR regions.

6. Steps d to g of claim 1 relate to the identification of Framework Region (FR) residues in the consensus human variable domain which are non-homologous to the corresponding import FR residues and replacement of such non-homologous human residues with corresponding import residues, if the residues are expected to have any one of the effects specified in step f. The locations of FR residues in human and mouse variable domains are indicated in Kabat *et al.* (see page ix) and the structural definition of the FR's was available (see Chothia *et al.*) Hence, it would have been straightforward for the skilled immunologist to identify the FR residues in the consensus human variable domain and the

import sequence. Using computer programs (such as the INSIGHT program [Biosym Technologies], available before June 14, 1991), the skilled biochemist would have been able to study the 3-dimensional structure of an antibody in order to establish whether a particular non-homologous import amino acid residue is likely to have one of the effects discussed in section f of claim 1. Information is provided on pages 14 to 16 of the specification which would have enabled the skilled biochemist to determine whether any non-homologous residue(s) would be expected to have the effects claimed. The techniques claimed in steps d to g of claim 1 could have been carried out routinely by a person versed in the relevant art, prior to June 14, 1991.

7. Steps a to g of claim 1 would have lead to the characterization of an amino acid sequence of a humanized antibody having non-human CDR amino acid residues and, optionally, having one or more non-human FR residues. In order to prepare the humanized antibody as claimed in claim 1, step h, the skilled biochemist could have synthesized the antibody using a peptide synthesizer which was commercially available before June 14, 1991. Alternatively, the antibody could have been made in recombinant cell culture (see page 26, last paragraph of the specification). Preparation of the antibody would have been straightforward to perform by the person skilled in the art, once the amino acid sequence of the humanized antibody had been characterized.

8. I understand that the Patent Office has rejected the above application on the basis that the sites in the variable domain referred to in claims 6, 7, and 9 are relevant to IgG antibodies only. It is my opinion that the sites referred to in claims 6, 7, and 9 would be relevant to other immunoglobulins. The basis for my opinion is given in paragraph 9 below.

9. The sites referred to in claims 6, 7, and 9 are the residue locations, or sites, of the FR residues in the heavy or light chain forming the variable domain of immunoglobulins. The residue sites referred to in claims 6, 7 & 9 relate to the position of a residue in the 3-D structure of the variable domain. Kabat *et al.* have used universal numbering for the amino acid residue locations of the variable domains for each of the immunoglobulin subgroups mentioned in the reference. The FR residue sites

indicated may be occupied by an amino acid residue which is non-homologous to the corresponding consensus human variable domain residue, and which is likely to have at least one of the effects discussed in step f of claim 1. These residue locations or sites are applicable *across species* (see page 16, line 8 of the specification). Accordingly, it is likely that an amino acid residue located at one of the sites indicated in claims 6, 7 and 9 will have one of the effects of claim 1 (step f), regardless of the antibody in which it is located, because it will be in the same position in the 3-D structure of the antibody variable domain as the residue sites referred to in the rejected claims. Accordingly, the examples of residue locations to be substituted in the variable domains would be applicable to antibodies, other than IgG antibodies.

10. I understand that the Patent Office has rejected the above application on the grounds that the invention as claimed is disclosed in Queen *et al.*, *Proc. Natl. Acad. Sci.*, **86**:10029-10033 (1989) or Co *et al.*, *Proc. Natl. Acad. Sci.*, **88**:2869-2873 (1991) and that the Office has suggested that the human variable domains disclosed in these references may have the same amino acid sequences as one of the consensus human variable domains disclosed in Kabat *et al.*

11. The above statements regarding the state of knowledge as of June 14, 1991, do not establish that the invention claimed in this application was known, or would have been obvious, to the skilled biochemist at the time the invention was made. To the contrary, after having read the citations relied upon by the Patent Office, it is my judgement that these documents would not have disclosed, nor suggested, the methods claimed. The basis for my opinion is given below.

12. The invention of the above application can be distinguished on the basis that a *consensus human variable domain* is used to "humanize" a non-human antibody of interest. The Queen *et al.* and Co *et al.* publications fail to disclose a consensus human variable domain. Instead, these publications refer to the use of a human variable domain having the closest sequence homology to the variable domain of the non-human antibody to be humanized. Queen *et al.* used the Eu human variable domain sequence (see Fig 2 thereof) and Co *et al.* used the variable domains of the Pom or Eu human

antibodies (see Fig 1 thereof). The sequences used in Queen *et al.* and Co *et al.* do not constitute a consensus human variable domain of a human immunoglobulin subgroup. The sequence identity between the amino acid sequences of the FR residues of the variable domains of the Pom or Eu heavy or light chains compared to the FR residues of the consensus human variable domains of each of the human immunoglobulin subgroups as defined by Kabat *et al.* is illustrated in Table 1 (see Exhibit "C", attached hereto). The CDR residues were not used in the comparison because of the large number of differences between these residues for variable domains of different antibodies. The Pom and Eu variable domain sequences were taken from Kabat *et al.* The consensus human variable domains of the V_L lambda subgroups IV and V were not compared, as these subgroups have too few members. While the variable domain of Eu is classified in subgroups V_L kappa I and V_H I, and the variable domain of Pom is classified in subgroups V_L kappa III and V_H III, it is apparent that the Eu and Pom variable domain amino acid sequences are not consensus human variable domains of any immunoglobulin subgroup. This is further demonstrated in the following paragraph.

13. Exhibits "D" and "C" attached hereto, show the differences in the amino acid sequences of the Pom and Eu heavy and light chain variable domains compared to the consensus human variable domain of the subgroup in which they are classified. Exhibit D illustrates an alignment of the amino acid sequences of the light chain variable domains of Eu, Pom and the consensus variable domain of the V_L kappa subgroup I (in which the light chain variable domain of Eu is classified). Exhibit E illustrates an alignment of the amino acid sequences of the heavy chain variable domains of Eu, Pom and the consensus variable domain of the V_H subgroup III (in which the heavy chain variable domain of Pom is classified). Even though Eu is classified in V_L kappa I, it has seven framework residues which are different from the framework residues of the kappa I consensus sequence. Furthermore, while Pom is classified in the V_H III subgroup, eight of its framework residues differ from the corresponding framework residues of the V_H III consensus sequence. There are, of course, many differences between the CDR residues of the consensus sequences and the corresponding CDR residues of Pom and Eu.

It is clear from the information in Exhibits C, D, & E that the Queen *et al.* and Co *et al.* publications fail to disclose a method wherein a non-human import antibody is humanized using a consensus human variable domain of an immunoglobulin subgroup.

14. I understand the Patent Office has rejected the above application on the basis that the invention claimed in claims 3 & 4 would have been obvious in light of Queen *et al.*, or Co *et al.*, when read in conjunction with Wallick *et al.*, *J. Exp. Med.*, **168** (1988). After reading these references, it is my opinion that the invention claimed in claims 3 and 4 is novel and would not have been obvious in light of the citations. The basis for my opinion is given in the following paragraph.

15. Claim 1 of the above application relates to a method of using a consensus human variable domain to "humanize" a non-human antibody (e.g. muMAb4D5). Use of a consensus human variable domain from a human immunoglobulin subgroup to humanize a non-human antibody is not disclosed in Queen *et al.*, Co *et al.* or Wallick *et al.* Wallick *et al.* does not relate to a method of humanizing a non-human antibody, much less a method of humanizing a non-human antibody using a consensus human variable domain of a human immunoglobulin subgroup. The skilled biochemist would have had no motivation at the filing date of this application to use a consensus human variable domain to humanize a non-human antibody, because the techniques in the prior literature had all relied upon using a human variable domain sequence which has the closest sequence homology to the non-human variable sequence (to be humanized) in order to reduce the likelihood of introducing distortions into the CDR's (see column 2 on page 10031 of Queen *et al.*) or to "retain high binding affinity in the humanized antibodies" (see column 1 on page 2871 of Co *et al.*). The method claimed in the above application does not rely on a high degree of homology between the variable domain of the non-human sequence and the consensus variable domain which is used to humanize the non-human sequence. It was surprising that a consensus variable domain of a selected immunoglobulin subgroup could be used to humanize a non-human antibody, regardless of the degree of homology between the human and non-human amino acid sequences. It was also surprising that the humanized antibody so formed retained,

and in some instances, had increased antigen binding affinity compared to the non-human antibody from which it was derived. The above application shows that the huMAb4D5-8 variant actually binds the p185^{HER2} ECD 3-fold more tightly than muMAb4D5 (see page 82 lines 31 & 32 to page 83, line 1 of the specification), which could not have been predicted by the ordinarily skilled biochemist at the time the specification was filed. Claim 3 refers to the step of finding any glycosylation site which is likely to affect the antigen binding or affinity in the import antibody and substituting the glycosylation site *into* the *consensus* amino acid sequence. Claim 4 refers to the step of *replacing* glycosylation sites of the consensus domain with the corresponding import amino acid residues if such glycosylation sites are not present in the import sequence. In my opinion, these claims would not have been obvious over the prior literature because the reference failed to disclose the use of a human consensus variable domain to humanize the non-human antibody. Accordingly, the skilled biochemist would have had no motivation to replace or insert glycosylation sites into a consensus amino acid sequence, as claimed in claims 3 and 4 of the application.

16. I declare further that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

Dated: 9/20/93

Signed: Robert F. Kelley
ROBERT F. KELLEY

CERTIFICATE OF MAILING

I hereby certify that this correspondence is being deposited with the United States Postal Service as first class mail in an envelope addressed to: Commissioner of Patents and Trademarks, Washington, D.C. 20231, on September 20, 1993.

Dated: September 20, 1993

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American Chemical Society, 1991-present

Scientific publications

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EXHIBIT C

TABLE 1
SEQUENCE IDENTITY - (%)

CONSENSUS VARIABLE DOMAIN SUBGROUP	EU	POM
V _L kappa I	92	76
V _L kappa II	61	71
V _L kappa III	72	85
V _L kappa IV	73	78
V _L lambda I	61	59
V _L lambda II	57	54
V _L lambda III	59	56
V _L lambda VI	52	49
V _H I	83	64
V _H II	53	62
V _H III	61	91

Variable Light Domain

	10	20	30	40
EU	DIQMTQSPSTLSASV	GDRVTITCRASQ	SINTWLAWYQOKPGKAPKLLMY	
		*	@ @ @	*
Kappa-I	DIQMTQSPSSLSASV	GDRVTITCRASQ	ISNYLAWYQOKPGKAPKLLIY	
	**	** * * * *	@ @ @ @	*** *
POM	EIVMTQSPVITLSVSPGERATL	SCRASQ	SISNYLAWYQOKPSGSPRLLIY	

CDR-L1

	50	60	70	80	90	100
EU	KASSLES	GVPSRFIGSGSGTEFTLT	ISSLOPDDFATYYCQYNSDKMFGQ			
	@	*	*	*	@ @ @	
Kappa-I	AASSLES	GVPSRFIGSGSGTEFTLT	ISSLOPDDFATYYCQYNSLPWTFGQ			
	@ @ @ @	* *	*	*	@ @ @	
POM	GASTRATGIPARFSGSGSGTEFTLT	ISSLOQSEDFAVYYCQYNNWPPTFGQ				

CDR-L2

CDR-L3

EU	GTKVEVKGT
	* *
Kappa-I	GTKVEIKRT
	*
POM	GTRVEIKR

KEY: * = differences in FR residues

@ = differences in CDR residues

EXHIBIT E

Variable Heavy Domain

	10	20	30	40
EU	QVQLVQSGAEVKKPGSSVKV	SCKASGGTFSRS	AIWVRQAPGGGLEWMG	
	* * * * *	* * * * *	@ @ @ @	* * *
human-III	EVQLVESGGGLVQPGGSLRL	SCAASGFTFSSY	AMSWVRQAPGKGLEWVS	
	*		@	*
POM	EVQLLESGGGLVQPGGSLRL	SCAASGFTFSSS	AMSWVRQAPGKGLEWVA	

CDR-H1

	50	a	60	70	80	abc	90
EU	GIVPMFGPPNYAQKFQGRVT	ITADESTNTAYMELSSLR	SEDTAFYFCAG				
	@ @ @ @ @	@ @ @ @	* * * * *	* * * * *			
human-III	VISGDGGSTIYADSVKGRFT	ISRDN	SKNTLYLQMN	SLRAEDTAVYYCAR			
	@ @ @ @ @	@	* * *	* * *			
POM	WKYENGNDKHYADSVNGRFT	ISRND	SKNTLYLLMNSL	QAEDTALYYCAR			

CDR-H2

		110
EU	GYGIYSPE----	EYNGGLVTVSS
	@ @ @ @ @	* * *
human-III	GRGGGSY----	WGQGTILTVSS
	@ @ @ @ @	*
POM	DAGPYVSPTFFAHYGQGT	LVT

CDR-H3

KEY: * = differences in FR residues

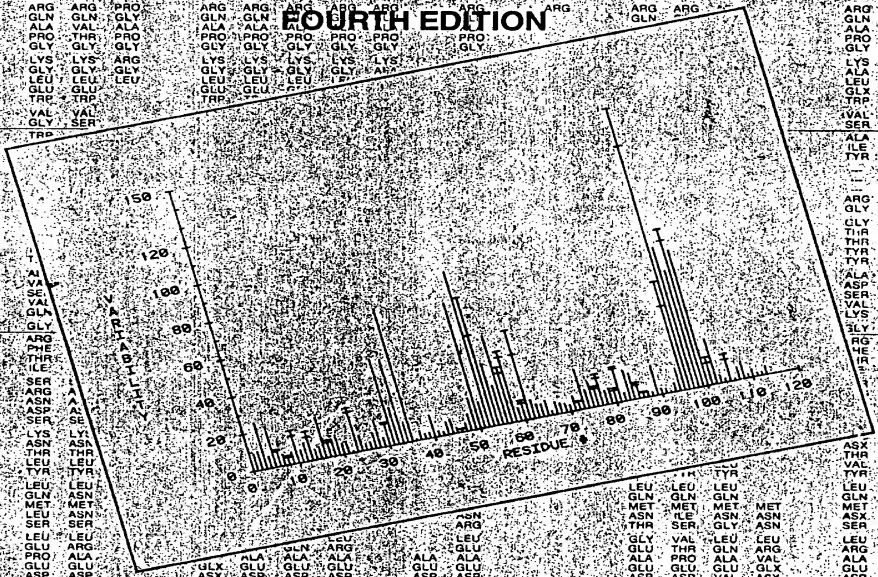
@ = differences in CDR residues

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SEQUENCES OF PROTEINS OF IMMUNOLOGICAL INTEREST

FOURTH EDITION



1987

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
Public Health Service
National Institutes of Health

SEQUENCES OF PROTEINS OF IMMUNOLOGICAL INTEREST

FOURTH EDITION

Tabulation and Analysis of
Amino Acid and Nucleic Acid Sequences of
Precursors, V-Regions, C-Regions, J-Chain,
T-Cell Receptor for Antigen, T-Cell Surface Antigens,
 β_2 -Microglobulins, Major Histocompatibility Antigens,
Thy-1, Complement, C-Reactive Protein, Thymopoietin,
Post-gamma Globulin, and α_2 -Macroglobulin

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considered uncertain by the authors have not been included in the table. In some instances the symbol # is used to indicate that several amino acid residues were found in one position, and these residues are listed in the notes. The four columns at the end of each table give:

1. the number of residues sequenced at that position,
2. the number of different amino acids found at that position,
3. the number of times the most common amino acid occurred and that amino acid in parentheses, and
4. the variability.

Variability is calculated (11) as:

$$\text{Variability} = \frac{\text{Number of different amino acids occurring at a given position}}{\text{Frequency of the most common amino acid at that position}}$$

An invariant position would have a variability of one; if 20 amino acids occurred with equal frequency, the variability would be 20 divided by 0.05 equals 400. If, for example, four different amino acids Ser, Asp, Pro, and Thr occurred at a given position, and of 100 sequences available at that position, Ser occurred 80 times, the variability would be $4/0.8 = 5$. When any of the amino acid residues sequenced were not identified completely and are listed as Glx (or Asx), two values, separated by a colon, are given in the last three columns. The first value in each of these columns is calculated assuming that only one of the two possibilities, e.g., Glu or Gln (or Asp or Asn) occurred, while the second considers that both were present and maximizes variability. In the variability plots, the horizontal bars indicate the two values.

When two or more amino acids are most common and occur with equal frequency, they are tabulated as a note, and the symbol + is used in the next to last column. If no sequence data have been reported for any position, there are no entries in the last four columns. Variability is not calculated for insertions or if only a single sequence is known. When the translated sequence of a clone corresponds to a previously listed sequence of a plasmacytoma from which it was prepared, only one sequence is listed so that the variability computations are not affected, and a note is included.

If a given sequence is associated with any antibody activity, this is indicated by an asterisk alongside the protein heading, and the antibody specificities are given in a separate list with binding constants if available. The notes list the a-allotypes for the rabbit heavy chain V-region and the b-allotypes for the constant domain of the rabbit kappa light chain. A key reference to the sequence is given; generally the most recent reference since it is usually the most nearly complete, but often several references are included, especially when revisions of a sequence have been made. Notes are now of two types; general notes about a table indicated by the symbol #, and specific notes indicated by the sequence number.

Signal Sequences

The signal (precursor) amino acid sequences of immunoglobulin chains are listed in three tables: one for kappa light chains, one for lambda light chains, and one for heavy chains. They were obtained either by direct sequencing of signal proteins (12-14) or by translating nucleotide sequences from DNA clones. Signal segments range from 17-29 amino acid residues in length and are thus numbered from -29 to -1. Genomic DNA clones contain introns of varying length that interrupt the coding sequence of the precursor within the codon for position -4, and in rare cases for position -6. Thus, the L-gene encodes the leader peptide to position -4 and the 5' end of the V-gene codes for positions -4 to -1.

The signal amino acid sequences of the T-cell receptors for antigens, β_2 -microglobulins, major histocompatibility complex proteins, and complement components are listed in separate tables.

By conformational energy calculations, the core V_{H} hydrophobic Leu-Leu-Leu-Trp-Val-Leu-Leu-Leu (MOPC321, MOPC63) exists in an alpha helical conformation, terminated by chain reversal conformations in the four C-terminal residues Trp-Val-Pro-Gly; the four amino terminal residues are compatible with the alpha helix (15).

Variable Region Sequences

The variable regions (16) of immunoglobulins have been shown to contain hypervariable segments in their light (11,17-23) and heavy (22,24-27) chains, of which certain residues have been affinity labeled (28-41). Three hypervariable segments of light chain were delineated from a statistical examination

of sequences of human V_{κ} , human V_{λ} , and mouse V_{κ} light chains aligned for maximum homology (11,22). These and the three corresponding segments of the heavy chains (22,26,27) were hypothesized (11,22) to be the complementarity-determining regions or segments (CDR) containing the residues which make contact with various antigenic determinants, and this has been verified by X-ray diffraction studies at high resolution (42-67). The rest of the V-region constitutes the framework (11,22,66-68). It is convenient to identify the framework segments (FR1, FR2, FR3, and FR4) and the complementarity-determining segments (CDR1, CDR2, and CDR3) with the three CDRs separating the four FRs. The residue numbers for these segments are as follows:

Segment	Light Chain	Heavy Chain
FR1	1-23 (with an occasional residue at 0, and a deletion at 10 in V_{λ} chains)	1-30 (with an occasional residue at 0)
CDR1	24-34 (with possible insertions numbered as 27A,B,C,D,E,F)	31-35 (with possible insertions numbered as 35A,B)
FR2	35-49	36-49
CDR2	50-56	50-65 (with possible insertions numbered as 52A,B,C) ^a
FR3	57-88	66-94 (with possible insertions numbered as 82A,B,C)
CDR3	89-97 (with possible insertions numbered as 95A,B,C,D,E,F)	95-102 (with possible insertions numbered as 100A,B,C,D,E,F,G,H,I,J,K)
FR4	98-107 (with a possible insertion numbered as 106A)	103-113

^a In the rabbit, Mage *et al.* (69) consider position 65 in V_H to be in FR3, since it is allotype related.

In the tables of V-regions, the FR and CDR are separated by horizontal lines for convenience in reading. One mouse kappa light chain, MPC11, has an extra segment of 12 amino acid residues between position 1 and the signal sequence (70). Several chains have internal deletions.

In the tables, the V-genes for the light chains code to amino acid position 95, and the J-minigenes from position 97 to 107 for lambda and 108 for kappa light chains. Position 96 is usually the site of V-J joining by recombination and may be coded partly by the V-gene and partly by the J-minigene. Because the site of V-J recombination could occur at different positions within a codon, different amino acid residues may result at this position. We have changed the location of the inserted residues from 97A-F (2) to 95A-F, since it makes for better alignment by confining chains of different lengths to the V-gene region. In V_{κ} chains, J1 and J2 were used 5 to 10 times more frequently than J4 and J5 (71).

The V-genes for the heavy chains code up to amino acid position 94 and are followed by the D- and J-minigenes. Because of the extensive variation in the lengths of D-minigenes, the exact boundary between D and J is not always located at the same amino acid position. In addition, the lengths of the J encoded amino acid sequences vary by a few amino acid residues. Moreover, the process of D-J joining appears to involve insertions of extra nucleotides between V and D and between D and J, termed the N region (72-76) and correlates with the appearance of terminal deoxynucleotidyl transferase in B cells (75). The original numbering system for the heavy chains has therefore been retained. Wysocki *et al.* (76) have provided some evidence suggesting a non-random origin for the V_H -D_H junction, perhaps a minigene, rather than random addition of the N nucleotides.

It has become evident that a critical understanding of the architecture of antibody combining sites and the genetics of the generation of diversity and of antibody complementarity will depend to a great extent on the evaluation of a large number of sequences of the variable regions and especially of the complementarity-determining segments of light and heavy chains of immunoglobulins of different species. Ability to locate residues in the site making contact with antigenic determinants (77) and to predict (67,78-82) the structures of antibody combining sites will depend heavily upon such sequences.

Figures 1 and 2 are stereoviews of the α -carbon skeletons of the four Fv regions for which high resolution X-ray structures have been determined, NEWM (44), KOL (62), MCP603 (47, 48, 63), and J539 (64). The residues in the CDRs are shown as solid circles. In Fig. 1 the combining site is at the

[illegible]

[illegible]

	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72
	HB	FR	GR	PAUL	MON	HEI	POT	S-GUI	SMYLOD	BL	SEB	JBL	PAO	CAR	MEV	BI	AMYL	CRA	DAV	FIN	KA	VL	LUX	NE	VA
	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
	0	1	2	3	4	5	6	7	8	9	10	11	12												

[illegible]

	97	98	99	100	101	102	103	104	105	106	107	108	109	110	111	# OF	# OF	OCCURRENCES
	LOD	HBJ	BEN	GR	MAA	MUK	AMY	AMR	AMC	BJ	HBJ	PEN	AMC	CL	CL	SE	AMINO	OF MOST COMMON
	10	10					554				6		MS					AMINO ACID
0	ASP	ASP	ASP	ASP	ASP	ASP	ASP	ASP	ASP	ASP	ASP	ASP	ASP	ASP	ASP	108	1	1(PCA)
1	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	107	3	103(ASP)
2	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	106	4	101(ILE)
3	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	105	8	97(GLN)
4	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	104	2	106(THR)
5	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	103	1	107(GLN)
6	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	102	3	100(SER)
7	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	101	4	105(SER)
8	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	100	5	98(SER)
9	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	99	7	91(THR)
10	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	98	4	91(ILE)
11	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	97	7	98(SER)
12	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	96	4	91(ILE)
13	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	95	7	88(SER)
14	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	94	3	91(ILE)
15	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	93	3	88(SER)
16	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	92	3	93(VAL)
17	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	91	3	90(ILE)
18	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	90	3	87(ASP)
19	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	89	6	88(VAL)
20	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	GLN	88	4	87(THR)
21	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	MET	87	4	84(ILE)
22	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	86	7	79(THR)
23	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	PRO	85	3	83(CYS)
24	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	SER	84	5	43(ARG)
25	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	ALA	83	4	71(ALA)
26	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	VAL	82	4	67(VAL)
27	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	THR	81	4	66(GLN)
28	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	ILE	80	3	53(GLN)
29	GLN	GLN	GLN															

HUMAN KAPPA LIGHT CHAINS SUBGROUP 1 (cont'd)
VARIABILITY

	0	
	1	3.2 : 4.4
	2	4.2
	3	6.9 : 9.3
	4	4.8
	5	3.1
	8	1. : 2.1
	7	3.2
	8	3.1
	9	4.2
F R 1	10	8.4
	11	5.7
	12	4.2
	13	4.4
	14	7.9
	15	3.1
	16	2. : 4.7
	17	3.2
	18	6.6
	19	3.1
	20	4.2
	21	4.2
	22	6.2
	23	1.
C D R 1	24	8.7
	25	4.2
	26	4.3
	27	4.4 : 5.4
	27A	
	27B	
	27C	
	27D	
	27E	
	27F	23. : 28.
	28	
	29	5.8
	30	19. : 21.
	31	26.
	32	14.
	33	4.3
	34	18. : 22.
F R 2	35	1.
	36	2.1
	37	4.3 : 4.9
	38	4.2 : 4.8
	39	4.4
	40	4.2
	41	3.3
	42	6.6
	43	2.2
	44	1.
	45	6.9
	46	9.8
	47	2.
	48	2.
	49	4.3
C D R 2	50	21. : 24.
	51	5.8
	52	4.3
	53	12. : 14.
	54	2.
	55	15.
	56	11.
	57	1.
	58	2.
	59	4.3
	60	1.
	61	3.1
F R 3	62	3.1
	63	8.4
	64	1.
	65	4.4
	66	3.1
	67	3.2
	68	3.2
	69	3.2
	70	8.2 : 11.
	71	4.3
	72	3.9
	73	4.3
	74	4.3
	75	3.2
	76	2.1
	77	7.4
	78	2.3
	79	2.1 : 3.4
	80	3.8
	81	4.1 : 7.7
	82	1. : 2.2
	83	3.7
	84	2.1
	85	4.3
	86	2.
	87	2.1
	88	1.
C D R 3	89	3.2 : 4.6
	90	3.3 : 5.1
	91	19. : 21.
	92	25. : 28.
	93	21.
	94	38.
	95	6.4
	95A	
	95B	
	95C	
	95D	
	95E	
F R 4	95F	43.
	96	4.6
	97	3.2
	98	1.
	99	6.9 : 9.1
	100	1.
	101	2.1
	102	1.
	103	5.4
	104	2.6
	105	4.5 : 6.3
	106	14.
	106A	
	107	2.1
	108	2.2
	109	1.

ANTIBODY SPECIFICITIES: HUMAN KAPPA LIGHT CHAINS SUBGROUP I

- 1) WEA: ANTI-3,4-PYRUVYLATED GALACTOSE MONOCLONAL
- 25) LOW: COLD AGGLUTININ WITH ANTI-BLOOD GROUP I ACTIVITY
- 39) LAY: ANTI-HUMAN GAMMA G1 AND G3 GLOBULINS: PO IDIOTYPE
- 51) HEI: COLD AGGLUTININ WITH ANTI-GD (MEMBRANE-GLYCOPOLIPID-DEPENDENT) ACTIVITY
- 66) DAY: ANTI-HUMAN GAMMA G GLOBULIN
- 67) FIN: ANTI-HUMAN GAMMA G GLOBULIN
- 72) WAG: ANTI-DINITROPHENYL
- 104) MAR: ANTI-LIPOPROTEIN LIPASE

ALLOTYPES: HUMAN KAPPA LIGHT CHAINS SUBGROUP I

- 79) KUE: INV(2)

CLASS: HUMAN KAPPA LIGHT CHAINS SUBGROUP I

- 8) WEA: IGM-KAPPA
- 33) F-GUI: IGG3-KAPPA
- 55) S-GUI: IGG3-KAPPA
- 74) PW: IGG1-KAPPA
- 82) RI: IGG1-KAPPA

REFERENCE: HUMAN KAPPA LIGHT CHAINS SUBGROUP I

- 1) ROY: HILSCHMANN N. & CRAIG L.C. (1965) PROC.NAT.ACAD.SCI.USA 53,1403-1409; HILSCHMANN N. (1967) Z.PHYSIOL.CHEM. 348,1077-1080; HILSCHMANN N. & BARRICELLI H. (1968) M.LANGE. & STENNETZKY H. (1969) PROC. 5TH FEBS SYMP. 15,57-74. (CHECKED BY AUTHOR WHO PROVIDED ADDITIONAL RESIDUES TO THOSE PUBLISHED AND CORRECTED RESIDUES 65 AND 87 AS GIVEN IN THE TABLE)
- 2) AU: SCHIECHL H. & HILSCHMANN N. (1971) Z.PHYSIOL.CHEM. 352,111-115; (1972) Z.PHYSIOL.CHEM. 353,345-370. (CHECKED BY AUTHOR)
- 3) REI: PALM W. & HILSCHMANN N. (1973) Z.PHYSIOL.CHEM. 354,1651-1654; (1976) Z.PHYSIOL.CHEM. 356,167-191. (CHECKED BY AUTHOR)
- 4) HAU: WATANABE S. & HILSCHMANN N. (1970) Z.PHYSIOL.CHEM. 351,1291-1295. (CHECKED BY AUTHOR)
- 5) HK101/CL: BENTLEY D.L. & RABBITS T.H. (1980) NATURE 288,730-733. (CHECKED BY AUTHOR 11/30/82)
- 6) SCW: EULITZ M., GYOTZ D. & HILSCHMANN N. (1972) Z.PHYSIOL.CHEM. 353,487-491; EULITZ M. & HILSCHMANN N. (1974) Z.PHYSIOL.CHEM. 355,842-866. (CHECKED BY AUTHOR)
- 7) AG: TITANI K., SHINDO A.T. & PUTNAM F.W. (1969) J.BIOL.CHEM. 244,3550-3560. (CHECKED BY AUTHOR 06/15/83)
- 8) WEA: GON P. & FRANGIONE B. (1963) PROC.NAT.ACAD.SCI.USA 40,4837-4841. (CHECKED BY AUTHOR 03/23/84)
- 9) HK137/CL: BENTLEY D.L. & RABBITS T.H. (1983) CELL 32,181-189.
- 10) HK134/CL: BENTLEY D.L. & RABBITS T.H. (1983) CELL 32,181-169.
- 11) DAUD/CL: KLOBECK H.G., COMBRATO G. & ZACHAU H.G. (1984) NUC.ACIDS RES. 12,18,6995-7006.
- 12) WALKER/CL: KLOBECK H.G., COMBRATO G. & ZACHAU H.G. (1984) NUC.ACIDS RES. 12,18,6995-7006. (CHECKED BY AUTHOR 08/22/85 WHO CORRECTED RESIDUE 34)
- 13) HF3-16/8: ATKINSON P.M., LAMPANANG W., FURIE B.C., NAPARSTEK Y., SCHWARTZ R.S., STOLLAR B.D. & FURIE B. (1985) J.CLIN.INVEST. 75,1138-1143. (CHECKED BY AUTHOR 08/21/85)
- 14) HF2-11/38: ATKINSON P.M., LAMPANANG W., FURIE B.C., NAPARSTEK Y., SCHWARTZ R.S., STOLLAR B.D. & FURIE B. (1985) J.CLIN.INVEST. 75,1138-1143. (CHECKED BY AUTHOR 08/21/85)
- 15) HF2-18/2: ATKINSON P.M., LAMPANANG W., FURIE B.C., NAPARSTEK Y., SCHWARTZ R.S., STOLLAR B.D. & FURIE B. (1985) J.CLIN.INVEST. 75,1138-1143. (CHECKED BY AUTHOR 08/21/85)
- 16) HF2-1/17: ATKINSON P.M., LAMPANANG W., FURIE B.C., NAPARSTEK Y., SCHWARTZ R.S., STOLLAR B.D. & FURIE B. (1985) J.CLIN.INVEST. 75,1138-1143. (CHECKED BY AUTHOR 08/21/85)
- 17) BJ26: ALESCIO-ZONTALI & BAGLIONI C. (1970) EUR.J.BIOCHEM. 15,450-463. (CHECKED BY AUTHOR)
- 18) RFZ: SMITHIES O., GIBSON D., FANNING E.M., GOODFLESH R.M., GILMAN J.G. & BALLANTYNE D.L. (1971) BIOCHEMISTRY 10,4912-4921. (CHECKED BY AUTHOR)
- 19) PSM: SEON B.K. (1982) MOL.IMMUNOL. 19,83-86. (CHECKED BY AUTHOR 05/23/83)
- 20) HOM: CAVIDIUD G., KLEIN M., HORNE C., HOFMANN T. & DORRINGTON K.J. (1981) MOL.IMMUNOL. 18,793-805.
- 21) ESM IGG: KUHN T.K., TUNG E., WANG I.Y. & WANG A.C. (1981) IMMUNOL. 44,265-271. (CHECKED BY AUTHOR 05/26/83)
- 22) ESM IGM: KUHN T.K., TUNG E., WANG I.Y. & WANG A.C. (1981) IMMUNOL. 44,265-271. (CHECKED BY AUTHOR 05/26/83)
- 23) WAT: STEVENSON J., WINTHOLM M.F.A., PANAGIOTPOULOS N., SCHIFFER M., POPP R.A. & SOLOMON A. (1981) J.MOL.BIOL. 147,185-193. (CHECKED BY AUTHOR 05/26/83)
- 24) AMYLOID VII-B: GLENNER G.G., TERRY W., HERADA M., ISERSKY C. & PAGE D. (1971) SCIENCE 172,1150-1151. (CHECKED BY AUTHOR 09/22/78)
- 25) LOW: CAPRA J.D., KEHOE J.M., WILLIAMS R.C., JR., FEIZI T. & KUNKEL H.G. (1970) PROC.NAT.ACAD.SCI.USA 69,40-43. (CHECKED BY AUTHOR WHO CORRECTED RESIDUE 18 AS GIVEN IN TABLE)
- 26) DIE: CAPRA J.D. & KUNKEL H.G. (1970) PROC.NAT.ACAD.SCI.USA 67,87-92. (CHECKED BY AUTHOR)
- 27) CAR A: CAPRA J.D. & KUNKEL H.G. (1970) PROC.NAT.ACAD.SCI.USA 67,87-92. (CHECKED BY AUTHOR)
- 28) TEI: CAPRA J.D. & KUNKEL H.G. (1970) PROC.NAT.ACAD.SCI.USA 67,87-92. (CHECKED BY AUTHOR)
- 29) BJ48: ALESCIO-ZONTALI & BAGLIONI C. (1970) EUR.J.BIOCHEM. 15,450-463. (CHECKED BY AUTHOR)
- 30) CON: NALL H.D. & EDMAN P. (1967) NATURE 216,262-263. (CHECKED BY AUTHOR 07/25/79)
- 31) TRA: NALL H.D. & EDMAN P. (1967) NATURE 216,262-263. (CHECKED BY AUTHOR 07/25/79)
- 32) AMYLOID LEP: LIAN J.B., SKINNER M., BENSON M.D. & COHEN A.S. (1977) BIOCHIM.BIOPHYS. ACTA 491,167-176.
- 33) F-GUI: WANG A.C., FUDENBERG H.H. & CREYSEL R. (1982) ACTA HAEMAT. 58,187-195. (CHECKED BY AUTHOR 05/26/83)
- 34) OUI/O: KOHLER H., SHIMIZU A., PAUL C. & PUTNAM F.W. (1970) SCIENCE 169,56-59; (KAPLAN A. & METZGER H. (1969) BIOCHEMISTRY 8,394-3951). (CHECKED BY AUTHOR 06/15/83)
- 35) DEE: MILSTEIN C. & DEVERSON E.V. (1971) BIOCHEM.J. 123,945-956. (CHECKED BY AUTHOR)
- 36) QAL/I: LAURE C.J., WATANABE S. & HILSCHMANN N. (1973) Z.PHYSIOL.CHEM. 354,1503-1504. (CHECKED BY AUTHOR)
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- 40) BRA: WANG A.C., WELLS J.V., FUDENBERG H.H. & GERGELY J. (1974) IMMUNOCHEM. 11,341-345. (CHECKED BY AUTHOR)
- 41) WEE: KRATZINH. J., KRUSCHE J.U. & HILSCHMANN N. (1980) Z.PHYSIOL.CHEM. 361,1591-1598.
- 42) VB/CL: PECH M., JAENICHEN H.-R., POHLNENZ H.-D., NEUMAIER P.S., KLOBECK H.-G. & ZACHAU H.G. (1984) J.MOL.BIOL. 176,189-204. (CHECKED BY AUTHOR 12/14/84)
- 43) VB/CL: PECH M., JAENICHEN H.-R., POHLNENZ H.-D., NEUMAIER P.S., KLOBECK H.-G. & ZACHAU H.G. (1984) J.MOL.BIOL. 176,189-204. (CHECKED BY AUTHOR 12/14/84)
- 44) HK102/CL: BENTLEY D.L. & RABBITS T.H. (1980) NATURE 288,730-733. (CHECKED BY AUTHOR 11/30/82)
- 45) EU: GOTTLEB D., CUNNINGHAM B.A., RUTISHAUSER U. & EDELMAN G.M. (1970) BIOCHEMISTRY 9,315-3161. (CHECKED BY AUTHOR)
- 46) DEN: YANG G.Y., PAULY E., KRATZINH. J. & HILSCHMANN N. (1981) Z.PHYSIOL.CHEM. 362,1131-1146.
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- 48) HBA: SMITH G.P., HOOD L. & FITCH W.M. (1971) ANN.REV.BIOCHEM. 40,969-1012.
- 49) FRA: MEINKEN G.C., SPIEGELBERG H.L. (1974) IMMUNOCHEM. 11,457-460. (CHECKED BY AUTHOR WHO PROVIDED ADDITIONAL RESIDUES TO THOSE PUBLISHED); MEINKEN G.C. & SPIEGELBERG H.L. (1978) IMMUNOCHEM. 13,975-979. (CHECKED BY AUTHOR 10/17/77)
- 50) GR: FAIR D.S., SLEDGE C., KRUEGER R.G., MANN K.G. & HOOD L.E. (1975) BIOCHEMISTRY 14,5581-5588.
- 51) PAU: SMITH G.P., HOOD L. & FITCH W.M. (1971) ANN.REV.BIOCHEM. 40,969-1012.
- 52) MON: NALL H.D. & EDMAN P. (1967) NATURE 216,262-263. (CHECKED BY AUTHOR 07/25/79)
- 53) HEI: RIESEN V.F., MAJANIEMI J., HUSER H., BRAUN D. & ROELCKE D. (1978) SCAND.J.IMMUNOL. 8,145-148. (CHECKED BY AUTHOR 10/10/79)
- 54) POT: CAPRA J.D. & KUNKEL H.G. (1970) PROC.NAT.ACAD.SCI.USA 67,87-92. (CHECKED BY AUTHOR WHO CORRECTED RESIDUE 9 AS GIVEN IN TABLE)
- 55) S-GUI: WANG A.C., FUDENBERG H.H. & CREYSEL R. (1982) ACTA HAEMAT. 58,187-195. (CHECKED BY AUTHOR 05/26/83)
- 56) AMYLOID BAN DWULET F.E., O'CONNOR T.P. & BENSON M.D. (1986) MOL.IMMUNOL. 23,73-78.
- 57) BJ19: ALESCIO-ZONTALI & BAGLIONI C. (1970) EUR.J.BIOCHEM. 15,450-463. (CHECKED BY AUTHOR)
- 58) BEL: MILSTEIN C. (1969) PROC. 5TH FEBS SYMP. 15,43-56. (CHECKED BY AUTHOR WHO PROVIDED ADDITIONAL RESIDUES TO THOSE PUBLISHED AND CORRECTED RESIDUES 1,3,6,27,79 AND 82 AS GIVEN IN TABLE)
- 59) JBL: SEON B.K. (1982) MOL.IMMUNOL. 19,83-86. (CHECKED BY AUTHOR 05/23/83)
- 60) PAP: NALL H.D. & EDMAN P. (1967) NATURE 216,262-263. (CHECKED BY AUTHOR 07/25/79)
- 61) CAR: MILSTEIN C.P. & DEVERSON E.V. (1974) EUR.J.BIOCHEM. 49,377-391. (CHECKED BY AUTHOR)
- 62) MEV: EULITZ M. & LINKE R.P. (1982) Z.PHYSIOL.CHEM. 363,1347-1358. (CHECKED BY AUTHOR 10/10/83)

NOTES: HUMAN KAPPA LIGHT CHAINS SUBGROUP I (cont'd)

GENERAL NOTES:

SEE SIGNAL PEPTIDE TABLE IF # OCCURS AT POSITION 0.

SPECIFIC NOTES:

- 5) HK101'CL: THE SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN FOETAL LIVER DNA.
- 7) AG: THE AMINO ACID RESIDUES AT POSITIONS 39 AND 41 WERE REPORTED BY THE AUTHORS AS GLY AND LYS RESPECTIVELY; HOWEVER, THE PROOF WAS NOT ABSOLUTE. THUS, THEY ARE OMITTED.
- 9) HK137'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN FETAL DNA.
- 10) HK134'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN FETAL DNA.
- 17) BJ28: ACID RESIDUES AT POSITIONS 39 AND 41 OF BJ28 WERE REPORTED BY THE AUTHORS AS GLY AND LYS RESPECTIVELY. SINCE THIS PROTEIN WAS SEQUENCED BEFORE THE SEQUENCES OF MANY OTHER PROTEINS WERE KNOWN AT THESE TWO POSITIONS, WE HAVE OMITTED THEM.
- 33) F-GUI: THE SEQUENCES OF F-GUI AND S-GUI WERE FROM THE SAME PATIENT.
- 44) HK102'CL: THE SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN FOETAL LIVER DNA.
- 55) S-GUI: THE SEQUENCES OF F-GUI AND S-GUI WERE FROM THE SAME PATIENT.
- 56) AMYLOID BAN: AMINO ACID RESIDUES FOUND AT POSITIONS 104 AND 105 ARE VALLEU AND GLN, GLU RESPECTIVELY.
- 57) BJ19: THE AMINO ACID RESIDUES AT POSITIONS 39 AND 41 WERE REPORTED BY THE AUTHORS AS GLY AND LYS RESPECTIVELY. SINCE THIS PROTEIN WAS SEQUENCED BEFORE THE SEQUENCES OF MANY OTHER PROTEINS WERE KNOWN AT THESE TWO POSITIONS, WE HAVE OMITTED THEM.
- 59) JBL: THE AMINO ACID RESIDUE FOUND AT POSITION 34 WAS ALA OR SER.
- 64) AMYLOID ES305: THE AMINO ACID RESIDUES AT POSITIONS 21 AND 29 WERE ILE OR LEU.
- 74) PW: THE SEQUENCE WAS FROM A PATIENT WITH TRANSITIONAL CELL CARCINOMA OF THE URINARY BLADDER.
- 82) RI: THE SEQUENCE WAS FROM A PATIENT WITH TRANSITIONAL CELL CARCINOMA OF THE URINARY BLADDER.
- 109) AMYLOID MS: THE AMINO ACID RESIDUE AT POSITION 2 MS WAS ILE OR LEU.
- 111) GM131'CL: FROM AN EPSTEIN-BARR VIRUS-TRANSFORMED HUMAN LYMPHOID CELL LINE

+ THE FOLLOWING WERE EQUALLY AND MOST FREQUENTLY OCCURRING:

AT POSITION	RESIDUES
27C	(LEU, VAL)
27D	(TRP, GLU)
50	(ALA, ASP)
92	(TYR, ASP, ASN)
95A	(SER, GLY)
95B	(TRP, GLY)

[illegible]

HUMAN KAPPA LIGHT CHAINS SUBGROUP II (cont'd)

	24*	25	26	27*	28	29	30	31	# OF	# OF	OCCURRENCES	VARIABILITY
	GIL	MEH	SC	TH	SYV	LUT	ROB	RAI	SEQUENCES	AMINO ACIDS	OF MOST COMMON AMINO ACID	
F R 1	0	ASP	ASP	ASP	ASP	ASP	ASP	ASP	31	1	31(ASP)	
	1	ILE	ILE	ILE	ILE	ILE	ILE	ILE	30	2	29(ILE)	1.
	2	VAL	VAL	VAL	VAL	VAL	VAL	VAL	30	2	28(VAL)	2.1
	3	MET	MET	MET	MET	MET	MET	MET	30	3	28(MET)	3.2
	4	THR	THR	THR	THR	THR	THR	THR	28	1	28(THR)	
	5	GLN	GLN	GLN	GLN	GLN	GLN	GLN	27	1	27(GLN)	1.
	6	SER	SER	SER	SER	SER	SER	SER	25	1	25(SER)	1.
	7	PRO	PRO	PRO	PRO	PRO	PRO	PRO	24	1	24(PRO)	1.
	8	LEU	LEU	LEU	LEU	LEU	LEU	LEU	25	1	25(LEU)	1.
	9	SER	SER	SER	SER	SER	SER	SER	24	1	24(SER)	1.
	10	LEU	LEU	LEU	LEU	LEU	LEU	LEU	24	1	24(LEU)	1.
	11	SER	SER	SER	SER	SER	SER	SER	24	1	23(SER)	2.1
	12	LEU	LEU	LEU	LEU	LEU	LEU	LEU	23	2	22(VAL)	2.1
	13	SER	SER	SER	SER	SER	SER	SER	17	1	17(THR)	1.
	14	THR	THR	THR	THR	THR	THR	THR	17	2	16(PRO)	2.1
	15	GLN	GLN	GLN	GLN	GLN	GLN	GLN	17	1	17(GLY)	1.
	16	THR	THR	THR	THR	THR	THR	THR	17	2	16(GLU)	2.1
	17	GLN	GLN	GLN	GLN	GLN	GLN	GLN	17	1	17(PRO)	1.
	18	THR	THR	THR	THR	THR	THR	THR	17	1	17(ALA)	1.
	19	GLN	GLN	GLN	GLN	GLN	GLN	GLN	17	1	17(SER)	1.
	20	THR	THR	THR	THR	THR	THR	THR	17	1	17(LEU)	1.
	21	GLN	GLN	GLN	GLN	GLN	GLN	GLN	17	2	16(SER)	2.1
	22	THR	THR	THR	THR	THR	THR	THR	17	1	17(CYS)	1.
C D R 1	23	ASP	ASP	ASP	ASP	ASP	ASP	ASP	16	1	16(ARG)	1.
	24	THR	THR	THR	THR	THR	THR	THR	14	2	13(SER)	2.2
	25	GLN	GLN	GLN	GLN	GLN	GLN	GLN	14	1	14(SER)	1.
	26	THR	THR	THR	THR	THR	THR	THR	14	1	14(GLN) 12(GLN)	1. 2.3
	27	GLN	GLN	GLN	GLN	GLN	GLN	GLN	12	3	10(SER)	
	28	THR	THR	THR	THR	THR	THR	THR	12	1	12(LEU)	
	29	GLN	GLN	GLN	GLN	GLN	GLN	GLN	12	3	9(LEU)	
	30	THR	THR	THR	THR	THR	THR	THR	10	5	8(HIS)	
	31	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	2	6(SER)	
	32	THR	THR	THR	THR	THR	THR	THR	7	2	7(ASP) 4(+) 4(-)	5.7 : 10.
	33	GLN	GLN	GLN	GLN	GLN	GLN	GLN	10	3	8(GLY)	3.8
	34	THR	THR	THR	THR	THR	THR	THR	9	4	3(ASN) 3(ASP) 2(+) 2(-)	9. : 12.
F R 2	35	GLN	GLN	GLN	GLN	GLN	GLN	GLN	9	1	9(TYR)	1.
	36	THR	THR	THR	THR	THR	THR	THR	8	1	8(LEU)	
	37	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	2	6(ASN) 4(+) 4(-)	2.7 : 4.
	38	THR	THR	THR	THR	THR	THR	THR	8	1	8(TRP)	
	39	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	2	7(TYR)	2.3
	40	THR	THR	THR	THR	THR	THR	THR	8	2	8(GLN) 6(GLN)	1. 2.7
	41	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	2	7(LYS)	2.3
	42	THR	THR	THR	THR	THR	THR	THR	8	1	7(PRO)	2.3
	43	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	1	8(GLY)	1.
	44	THR	THR	THR	THR	THR	THR	THR	8	1	8(GLN) 6(GLN)	1. 2.7
	45	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	1	6(SER)	
	46	THR	THR	THR	THR	THR	THR	THR	7	1	7(PRO)	
	47	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	2	5(GLN) 3(+) 3(-)	4.2 : 7.
C D R 2	48	THR	THR	THR	THR	THR	THR	THR	7	1	6(LEU)	2.3
	49	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	1	7(LEU)	1.
	50	THR	THR	THR	THR	THR	THR	THR	7	1	7(ILE)	1.
	51	GLN	GLN	GLN	GLN	GLN	GLN	GLN	6	1	6(TYR)	1.
	52	THR	THR	THR	THR	THR	THR	THR	6	4	4(LEU)	4.5
	53	GLN	GLN	GLN	GLN	GLN	GLN	GLN	6	1	3(GLY)	8.
	54	THR	THR	THR	THR	THR	THR	THR	7	2	7(SER)	1.
	55	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	2	5(ASN)	2.6
	56	THR	THR	THR	THR	THR	THR	THR	7	1	7(ARG)	1.
	57	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	2	7(SALA)	2.6
	58	THR	THR	THR	THR	THR	THR	THR	7	1	7(SED)	1.
	59	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	1	7(GLY)	1.
	60	THR	THR	THR	THR	THR	THR	THR	7	1	7(VAL)	1.
F R 3	61	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	1	7(TYR)	1.
	62	THR	THR	THR	THR	THR	THR	THR	7	1	7(ASP) 6(ASP)	1. 2.3
	63	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	1	7(ARG)	1.
	64	THR	THR	THR	THR	THR	THR	THR	8	1	8(PHE)	1.
	65	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	1	8(SER)	1.
	66	THR	THR	THR	THR	THR	THR	THR	8	2	7(GLY)	2.3
	67	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	1	8(SER)	1.
	68	THR	THR	THR	THR	THR	THR	THR	8	1	8(GLY)	1.
	69	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	1	8(SER)	1.
	70	THR	THR	THR	THR	THR	THR	THR	8	2	7(GLY)	2.3
	71	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	1	7(THR)	1.
	72	THR	THR	THR	THR	THR	THR	THR	7	1	7(ASP) 6(ASP)	1. 2.3
	73	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	1	8(PHE)	1.
C D R 3	74	THR	THR	THR	THR	THR	THR	THR	8	1	8(THR)	1.
	75	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	1	8(LEU)	1.
	76	THR	THR	THR	THR	THR	THR	THR	8	3	8(LYS)	4.
	77	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	3	8(LEU)	4.
	78	THR	THR	THR	THR	THR	THR	THR	8	2	7(SER)	2.3
	79	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	1	8(ARG)	1.
	80	THR	THR	THR	THR	THR	THR	THR	8	1	8(VAL)	1.
	81	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	2	8(GLU) 4(+) 4(-)	2.7 : 4.
	82	THR	THR	THR	THR	THR	THR	THR	8	1	7(ALA)	1.
	83	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	1	8(GLU) 6(GLU)	1. 2.7
	84	THR	THR	THR	THR	THR	THR	THR	8	1	8(ASP) 6(ASP)	1. 2.7
	85	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	1	8(GLY)	1.
	86	THR	THR	THR	THR	THR	THR	THR	8	1	8(VAL)	1.
F R 4	87	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	1	8(TYR)	1.
	88	THR	THR	THR	THR	THR	THR	THR	8	1	8(TYR)	1.
	89	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	1	8(CYS)	1.
	90	THR	THR	THR	THR	THR	THR	THR	7	1	7(MET)	1.
	91	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	1	7(GLN) 6(GLN)	1. 2.3
	92	THR	THR	THR	THR	THR	THR	THR	7	2	7(SALA)	4.2
	93	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	3	5(LEU)	2.6 : 5.3
	94	THR	THR	THR	THR	THR	THR	THR	7	5	5(GLN) 4(GLN)	4.2
	95	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	2	2(+) 2(-)	18.
	96	THR	THR	THR	THR	THR	THR	THR	7	2	6(PRO)	2.3
	97	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	6	2(TYR)	21.
	98	THR	THR	THR	THR	THR	THR	THR	7	1	7(THR)	1.
	99	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	1	7(PHE)	1.
F R 5	100	THR	THR	THR	THR	THR	THR	THR	7	1	7(GLY)	1.
	101	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	2	6(GLN)	2.3
	102	THR	THR	THR	THR	THR	THR	THR	7	1	7(GLY)	1.
	103	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	1	7(THR)	1.
	104	THR	THR	THR	THR	THR	THR	THR	7	2	5(LYS)	4.2
	105	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	1	4(+) 4(-)	4.
	106	THR	THR	THR	THR	THR	THR	THR	8	1	8(GLU) 6(GLU)	1. 2.3
	107	GLN	GLN	GLN	GLN	GLN	GLN	GLN	8	1	8(ILE)	1.
	108	THR	THR	THR	THR	THR	THR	THR	8	2	7(LYS)	2.3
	109	GLN	GLN	GLN	GLN	GLN	GLN	GLN	7	1	7(ARG)	1.
	110	THR	THR	THR	THR	THR	THR	THR	4	1	4(THR)	1.

ANTIBODY SPECIFICITIES: HUMAN KAPPA LIGHT CHAINS SUBGROUP II

- 3) ROB: COLD AGGLUTININ WITH ANTI-PRID ACTIVITY
 10) WILS: COLD AGGLUTININ WITH ANTI-BLOOD GROUP I ACTIVITY
 14) FR: ANTI-PHOSPHOCHOLINE(BINDING CONSTANT = 6.4X10⁶)
 24) NIL: ANTI-HGG
 27) TH: COLD AGGLUTININ WITH ANTI-PP2A ACTIVITY (RBC MEMBRANE ANTIGEN ON HUMAN, RAT AND GUINEA PIG ERYTHROCYTES INACTIVATED BY PROTEOLYTIC ENZYMES AND NEURAMINIDASE)

REFERENCE: HUMAN KAPPA LIGHT CHAINS SUBGROUP II

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 3) NIM: EULITZ M. & KLEY H.-P. (1977) IMMUNOCHEM. 14,289-297. (CHECKED BY AUTHOR 10/18/77)
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 5) GM 607: CL: KLOBECK H.G. SOLOMONA A. & ZACHAU H.G. (1984) NATURE 309,73-76.
 6) BAT: DAYHOFF M.O. (1972) ATLAS OF PROTEIN SEQUENCE & STRUCTURE 5.D-246. SUBMITTED BY SMITHES O., GIBSON D.M. AND FANNING E.M. (CHECKED BY AUTHOR)
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 8) ROB: GERGELY J., WANG A.C. & FUDENBERG H.H. (1973) VOX SANG. 24,432-440. (CHECKED BY AUTHOR)
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 11) QLI: FRANGIONE B., FRANKLIN E.C. & PRELLI F. (1976) SCAND. J. IMMUNOL. 5,623-627. (CHECKED BY AUTHOR 10/17/77)
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 16) RPM1-6410: CL: HIETTER P.A., MAX E.E., SEIDMAN J.O., MAIZEL J.V., JR. & LEDER P. (1980) CELL 22,197-207; KLOBECK H.G., MEINDL G., COMBIATO G., SOLOMONA A. & ZACHAU H.G. (1985) NUCLEIC ACIDS RES. 13, 849-856.
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 24) QLI: ABRAHAM G.N., BROWN P., JOHNSTON S.L., NELLIS L., MARKS S. & WELCH E.H. (1978) IMMUNOLOGY 35,447-453. (CHECKED BY AUTHOR 07/23/79)
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NOTES: HUMAN KAPPA LIGHT CHAINS SUBGROUP II

IDENTICAL SETS OF FRAMEWORK SEGMENTS:

- FR1: SET 1: TEW(1), MIL(2), NIM(3), CUM(4), GM 607 'CL(5), BAT(6), BATES(7), ROB(8), SLO(9), WILS(10), QLI(11), AMYLOID TWP(12), RAI(13). (13 IDENTICAL)
 FR2: SET 1: MIL(2), NIM(3), GM 607 'CL(5). (3 IDENTICAL HUMAN V-KAPPA-II; ALSO 2 MOUSE V-KAPPA-II: VKAPPA 24B'CL(63), 251(367))
 SET 2: MIL(2), FR(14). (2 IDENTICAL)
 FR3: SET 1: TEW(1), GM 607 'CL(5), RPM1-6410'CL(16). (3 IDENTICAL)
 FR4: SET 1: GM 607 'CL(5), RPM1-6410'CL(16). (2 IDENTICAL HUMAN V-KAPPA-II; ALSO 3 HUMAN V-KAPPA-I: AUJ(2), GAL(II)(36), CL(110); 7 HUMAN V-KAPPA-III: V(37), PAY(17), PIE(11), GLO(15), CUR(20), REE(15), VKAPPA3'CL(82); AND 1 HUMAN V-KAPPA-IV: PB17V'CL(31))
 SET 2: NIM(3), FR(14). (2 IDENTICAL HUMAN V-KAPPA-II; ALSO 3 HUMAN V-KAPPA-I: AG(7), DEN(46), B(83); 6 HUMAN V-KAPPA-III: NEU(5), GO(16), GAR(10), FLO(12), FR(12), IARC(BL4)'CL(28); AND 1 HUMAN V-KAPPA-IV: LEN(41))
 SET 3: TEW(1). (IDENTICAL TO 2 HUMAN V-KAPPA-I: WALKER'CL(12), OUI(10)(34))

IDENTICAL SETS OF COMPLEMENTARY TYPING REGIONS:

- CDR1:
 CDR2: SET 1: MIL(2), NIM(3), GM 607 'CL(5). (3 IDENTICAL)
 CDR3:

IDENTICAL SETS OF J-MINIGENES:

- SET 1: RPM1-6410'CL(16). (IDENTICAL TO 1 HUMAN V-KAPPA-I: AUJ(2); 2 HUMAN V-KAPPA-III: PIE(11), VKAPPA3'CL(82); AND 1 HUMAN V-KAPPA-IV: PB17V'CL(31))
 SET 2: TEW(1). (IDENTICAL TO 1 HUMAN V-KAPPA-I: WALKER'CL(12))
 SET 3: FR(14). (IDENTICAL TO 2 HUMAN V-KAPPA-I: DEN(46), B(83); AND 3 HUMAN V-KAPPA-III: GAR(10), FLO(12), IARC(BL4)'CL(28))

SPECIFIC NOTES:

- 12) AMYLOID TWP: IT HAS THE SAME SEQUENCE AS THAT OF TEW SO FAR AS THE SEQUENCED POSITIONS ARE CONCERNED.
 14) FR: AN IDIOTYPIC ANTIBODY TO FR NOT INHIBITED BY PHOSPHORYLCHOLINE REACTED BETTER WITH THE FR HEAVY CHAIN THAN WITH THE LIGHT CHAIN. THE CROSS-REACTION WITH MOPC167 WAS 10,000 TIMES WEAKER. (RIESEN W.F. (1979) EUR. J. IMMUNOL., 9,421-425).
 16) RPM1-6410:CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN ADULT DNA.

+ THE FOLLOWING WERE EQUALLY AND MOST FREQUENTLY OCCURRING:

AT POSITION	RESIDUES
27F	(GLY,ASN) (GLY,ASP)
28	(ASP,ASN)
31	(THR,ASP)
34	(ASP,ASP)
45	(GLU,GLN)
79	(GLU,GLN)
94	(THR,SER)
104	(LEU,VAL)

	IN	VAR	RES	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105	106	107	108	109	110	111	112	113	114	115	116	117	118	119	120	121	122	123	124	125	126	127	128	129	130	131	132	133	134	135	136	137	138	139	140	141	142	143	144	145	146	147	148	149	150	151	152	153	154	155	156	157	158	159	160	161	162	163	164	165	166	167	168	169	170	171	172	173	174	175	176	177	178	179	180	181	182	183	184	185	186	187	188	189	190	191	192	193	194	195	196	197	198	199	200	201	202	203	204	205	206	207	208	209	210	211	212	213	214	215	216	217	218	219	220	221	222	223	224	225	226	227	228	229	230	231	232	233	234	235	236	237	238	239	240	241	242	243	244	245	246	247	248	249	250	251	252	253	254	255	256	257	258	259	260	261	262	263	264	265	266	267	268	269	270	271	272	273	274	275	276	277	278	279	280	281	282	283	284	285	286	287	288	289	290	291	292	293	294	295	296	297	298	299	300	301	302	303	304	305	306	307	308	309	310	311	312	313	314	315	316	317	318	319	320	321	322	323	324	325	326	327	328	329	330	331	332	333	334	335	336	337	338	339	340	341	342	343	344	345	346	347	348	349	350	351	352	353	354	355	356	357	358	359	360	361	362	363	364	365	366	367	368	369	370	371	372	373	374	375	376	377	378	379	380	381	382	383	384	385	386	387	388	389	390	391	392	393	394	395	396	397	398	399	400	401	402	403	404	405	406	407	408	409	410	411	412	413	414	415	416	417	418	419	420	421	422	423	424	425	426	427	428	429	430	431	432	433	434	435	436	437	438	439	440	441	442	443	444	445	446	447	448	449	450	451	452	453	454	455	456	457	458	459	460	461	462	463	464	465	466	467	468	469	470	471	472	473	474	475	476	477	478	479	480	481	482	483	484	485	486	487	488	489	490	491	492	493	494	495	496	497	498	499	500	501	502	503	504	505	506	507	508	509	510	511	512	513	514	515	516	517	518	519	520	521	522	523	524	525	526	527	528	529	530	531	532	533	534	535	536	537	538	539	540	541	542	543	544	545	546	547	548	549	550	551	552	553	554	555	556	557	558	559	560	561	562	563	564	565	566	567	568	569	570	571	572	573	574	575	576	577	578	579	580	581	582	583	584	585	586	587	588	589	590	591	592	593	594	595	596	597	598	599	600	601	602	603	604	605	606	607	608	609	610	611	612	613	614	615	616	617	618	619	620	621	622	623	624	625	626	627	628	629	630	631	632	633	634	635	636	637	638	639	640	641	642	643	644	645	646	647	648	649	650	651	652	653	654	655	656	657	658	659	660	661	662	663	664	665	666	667	668	669	670	671	672	673	674	675	676	677	678	679	680	681	682	683	684	685	686	687	688	689	690	691	692	693	694	695	696	697	698	699	700	701	702	703	704	705	706	707	708	709	710	711	712	713	714	715	716	717	718	719	720	721	722	723	724	725	726	727	728	729	730	731	732	733	734	735	736	737	738	739	740	741	742	743	744	745	746	747	748	749	750	751	752	753	754	755	756	757	758	759	760	761	762	763	764	765	766	767	768	769	770	771	772	773	774	775	776	777	778	779	780	781	782	783	784	785	786	787	788	789	790	791	792	793	794	795	796	797	798	799	800	801	802	803	804	805	806	807	808	809	810	811	812	813	814	815	816	817	818	819	820	821	822	823	824	825	826	827	828	829	830	831	832	833	834	835	836	837	838	839	840	841	842	843	844	845	846	847	848	849	850	851	852	853	854	855	856	857	858	859	860	861	862	863	864	865	866	867	868	869	870	871	872	873	874	875	876	877	878	879	880	881	882	883	884	885	886	887	888	889	890	891	892	893	894	895	896	897	898	899	900	901	902	903	904	905	906	907	908	909	910	911	912	913	914	915	916	917	918	919	920	921	922	923	924	925	926	927	928	929	930	931	932	933	934	935	936	937	938	939	940	941	942	943	944	945	946	947	948	949	950	951	952	953	954	955	956	957	958	959	960	961	962	963	964	965	966	967	968	969	970	971	972	973	974	975	976	977	978	979	980	981	982	983	984	985	986	987	988	989	990	991	992	993	994	995	996	997	998	999	1000
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HUMAN KAPPA LIGHT CHAINS SUBGROUP III (cont'd)

	75 DOB	76 H56	77 H5J 12	78 BUR R	79 LEG	80 B6	81 AMYLOID WR	82 VK AP3	CL	# OF SEQUENCES	# OF AMINO ACIDS	OCCURRENCES OF MOST COMMON AMINO ACID	VARIABILITY
F R 1	0	---	---	---	---	---	---	---	---	79	3	74(GLU) 73(GLU)	3.2 : 4.3
	1	GLU	GLU	GLU	GLU	---	---	---	---	79	5	74(ILE)	5.3
	2	ILE	ILE	ILE	ILE	---	---	---	---	79	4	76(VAL)	4.2
	3	MEI	VAL	VAL	VAL	---	---	---	---	79	3	85(LEU)	3.8
	4	---	---	---	---	---	---	---	---	77	1	77(THR)	1.
	5	THR	THR	THR	THR	---	---	---	---	75	1	75(GLN) 69(GLN)	2.1 : 2.2
	6	GLN	GLN	GLN	---	---	---	---	---	75	1	75(SER)	1.
	7	PRO	---	---	---	---	---	---	---	74	1	74(PRO)	1.
	8	ALA	---	---	---	---	---	---	---	69	7	46(GLY)	9. : 11.
	9	---	---	---	---	---	---	---	---	70	4	68(THR)	4.2
	10	---	---	---	---	---	---	---	---	68	4	67(LEU)	2.
	11	---	---	---	---	---	---	---	---	67	1	67(SER)	1.
	12	---	---	---	---	---	---	---	---	67	1	62(LEU)	6.4
	13	---	---	---	---	---	---	---	---	66	2	64(SER)	2.1
C D 2	14	---	---	---	---	---	---	---	---	66	2	65(PRO)	2.
	15	---	---	---	---	---	---	---	---	62	3	62(GLY)	1.
	16	---	---	---	---	---	---	---	---	62	3	56(GLU) 50(GLU)	3.3 : 5.
	17	---	---	---	---	---	---	---	---	58	7	11(ARG)	8.
	18	---	---	---	---	---	---	---	---	60	2	52(ALA)	2.3
	19	---	---	---	---	---	---	---	---	59	5	53(THR)	5.6
	20	---	---	---	---	---	---	---	---	60	5	57(LEU)	2.1
	21	---	---	---	---	---	---	---	---	50	3	56(SER)	3.1
	22	---	---	---	---	---	---	---	---	50	3	50(CYS)	1.
	23	---	---	---	---	---	---	---	---	51	4	47(ARG)	4.3
	24	---	---	---	---	---	---	---	---	52	3	51(ALA)	2.
	25	---	---	---	---	---	---	---	---	49	3	46(SER)	1.
	26	---	---	---	---	---	---	---	---	47	3	43(GLN) 37(GLN)	3.3 : 3.8
	27A	---	---	---	---	---	---	---	---	32	4	25(SER)	1.
C D 3	27B	---	---	---	---	---	---	---	---	---	---	---	---
	27C	---	---	---	---	---	---	---	---	---	---	---	---
	27D	---	---	---	---	---	---	---	---	---	---	---	---
	27E	---	---	---	---	---	---	---	---	---	---	---	---
	27F	---	---	---	---	---	---	---	---	---	---	---	---
	28	---	---	---	---	---	---	---	---	47	7 : 8	25(VAL)	13. : 15.
	29	---	---	---	---	---	---	---	---	44	6	27(SER)	9.8
	30	---	---	---	---	---	---	---	---	40	7	24(SER)	12.
	31	---	---	---	---	---	---	---	---	39	10	24(SER)	16.
	32	---	---	---	---	---	---	---	---	40	8	26(TYR)	11.
	33	---	---	---	---	---	---	---	---	41	4	36(LEU)	4.6
	34	---	---	---	---	---	---	---	---	41	5	37(ALA)	5.5
	35	---	---	---	---	---	---	---	---	38	1	38(TRP)	1.
C D 4	36	---	---	---	---	---	---	---	---	39	1	39(TYR)	1.
	37	---	---	---	---	---	---	---	---	39	2	39(GLN) 33(GLN)	1. : 2.4
	38	---	---	---	---	---	---	---	---	37	2 : 3	36(GLN) 30(GLN)	2.1 : 3.7
	39	---	---	---	---	---	---	---	---	33	3	29(LYS)	3.
	40	---	---	---	---	---	---	---	---	34	3	32(PRO)	3.2
	41	---	---	---	---	---	---	---	---	27	4	26(GLY)	2.1
	42	---	---	---	---	---	---	---	---	27	4	24(GLN) 23(GLN)	4.5 : 4.7
	43	---	---	---	---	---	---	---	---	26	3	23(ALA)	3.4
	44	---	---	---	---	---	---	---	---	27	3	25(PRO)	3.2
	45	---	---	---	---	---	---	---	---	26	3	24(ARG)	3.3
	46	---	---	---	---	---	---	---	---	23	3	23(LEU)	3.1
	47	---	---	---	---	---	---	---	---	23	2	22(LEU)	2.1
	48	---	---	---	---	---	---	---	---	22	3	20(LE)	3.3
	49	---	---	---	---	---	---	---	---	22	4	19(TYR)	4.6
C D 5	50	---	---	---	---	---	---	---	---	21	5	16(GLY)	6.6
	51	---	---	---	---	---	---	---	---	20	2	16(LEU)	2.8
	52	---	---	---	---	---	---	---	---	20	2	18(SER)	2.2
	53	---	---	---	---	---	---	---	---	20	2	18(LEU)	2.6
	54	---	---	---	---	---	---	---	---	20	2	18(ARG)	2.1
	55	---	---	---	---	---	---	---	---	23	3	21(ALA)	3.3
	56	---	---	---	---	---	---	---	---	22	2	19(THR)	4.6
	57	---	---	---	---	---	---	---	---	23	2	22(GLY)	2.1
	58	---	---	---	---	---	---	---	---	23	3	21(ILE)	3.3
	59	---	---	---	---	---	---	---	---	23	1	23(PRO)	1.
	60	---	---	---	---	---	---	---	---	23	5	17(ASP)	6.8
	61	---	---	---	---	---	---	---	---	23	1	23(PHE)	1.
	62	---	---	---	---	---	---	---	---	23	1	21(SER)	2.2
	63	---	---	---	---	---	---	---	---	23	2	23(GLY)	1.
C D 6	64	---	---	---	---	---	---	---	---	23	1	21(SER)	2.1
	65	---	---	---	---	---	---	---	---	22	2	17(GLY)	5.2
	66	---	---	---	---	---	---	---	---	22	2	21(SER)	2.1
	67	---	---	---	---	---	---	---	---	22	1	22(GLY)	1.
	68	---	---	---	---	---	---	---	---	22	2	21(THR)	2.1
	69	---	---	---	---	---	---	---	---	22	2	21(THR)	2.1
	70	---	---	---	---	---	---	---	---	21	2	19(ASP)	2.2
	71	---	---	---	---	---	---	---	---	21	1	21(PHE)	1.
	72	---	---	---	---	---	---	---	---	21	1	21(THR)	1.
	73	---	---	---	---	---	---	---	---	21	1	21(LEU)	1.
	74	---	---	---	---	---	---	---	---	21	2	20(THR)	2.1
	75	---	---	---	---	---	---	---	---	21	3	20(LE)	3.2
	76	---	---	---	---	---	---	---	---	21	3	19(SER)	3.3
	77	---	---	---	---	---	---	---	---	22	5	16(ARG)	6.9
C D 7	78	---	---	---	---	---	---	---	---	22	5	20(LEU)	1.
	79	---	---	---	---	---	---	---	---	22	2	21(GLU) 20(GLU)	2.1 : 2.2
	80	---	---	---	---	---	---	---	---	22	2	19(PRO)	2.
	81	---	---	---	---	---	---	---	---	22	1	21(GLU)	2.1
	82	---	---	---	---	---	---	---	---	22	2	22(ASP)	2.1
	83	---	---	---	---	---	---	---	---	22	3	20(PHE)	3.3
	84	---	---	---	---	---	---	---	---	22	1	22(ALA)	1.
	85	---	---	---	---	---	---	---	---	22	1	21(VAL)	2.1
	86	---	---	---	---	---	---	---	---	22	1	22(TYR)	2.2
	87	---	---	---	---	---	---	---	---	22	2	20(TYR)	2.2
	88	---	---	---	---	---	---	---	---	22	1	22(CYS)	1.
	89	---	---	---	---	---	---	---	---	22	2	21(GLN)	2.1
	90	---	---	---	---	---	---	---	---	22	1	22(GLN)	1.
	91	---	---	---	---	---	---	---	---	22	2	20(TYR)	2.2
	92	---	---	---	---	---	---	---	---	22	2	16(GLY)	6.9
C D 8	93	---	---	---	---	---	---	---	---	21	5	12(SER)	8.8
	94	---	---	---	---	---	---	---	---	21	3	18(SER)	4.7
	95	---	---	---	---	---	---	---	---	21	1	18(PRO)	3.5
	95A	---	---	---	---	---	---	---	---	1	1	1(PRO)	1.
	95B	---	---	---	---	---	---	---	---	---	---	---	---
	95C	---	---	---	---	---	---	---	---	---	---	---	---
	95D	---	---	---	---	---	---	---	---	---	---	---	---
	95E	---	---	---	---	---	---	---	---	---	---	---	---
	96	---	---	---	---	---	---	---	---	19	10	4(TYR)	48.
	97	---	---	---	---	---	---	---	---	20	1	19(THR)	2.1
	98	---	---	---	---	---	---	---	---	20	1	20(PHE)	1.
	99	---	---	---	---	---	---	---	---	20	1	20(GLY)	1.
	100	---	---	---	---	---	---	---	---	20	2	18(GLN)	2.2
	101	---	---	---	---	---	---	---	---	20	1	20(GLY)	2.
	102	---	---	---	---	---	---	---	---	20	2	18(THR)	2.2
C D 9	103	---	---	---	---	---	---	---	---	20	2	18(LYS)	2.2
	104	---	---	---	---	---	---	---	---	20	2	11(VAL)	2.2
	105	---	---	---	---	---	---	---	---	20	3	18(GLU)	2.2
	106	---	---	---	---	---	---	---	---	20	3	18(ILE)	2.2
	106A	---	---	---	---	---	---	---	---	---	---	---	---
	107	---	---	---	---	---	---	---	---	20	2	18(LYS)	2.1
	108	---	---	---	---	---	---	---	---	16	1	18(ARG)	1.
	109	---	---	---	---	---	---	---	---	10	1	10(THR)	1.

ANTIBODY SPECIFICITIES: HUMAN KAPPA LIGHT CHAINS SUBGROUP III

- 2) WOL: ANTI-HUMAN GAMMA G GLOBULIN; WA IDIOTYPE
- 3) SIE: ANTI-HUMAN GAMMA G GLOBULIN; WA IDIOTYPE
- 5) NEU: CRYOGLOBULIN WITH ANTI-HGG ACTIVITY; B IDIOTYPE (KUNKEL H.G. WINCHESTER R.J. JOSLIN F.G. & CAPRA J.D. (1974) J EXP MED. 139, 128)
- 6) GOT: CRYOGLOBULIN WITH ANTI-HGG ACTIVITY; B IDIOTYPE
- 7) PAY: CRYOGLOBULIN WITH ANTI-HGG ACTIVITY; B IDIOTYPE
- 8) SON: CRYOGLOBULIN WITH ANTI-LOW-DENSITY LIPOPROTEIN ACTIVITY; B IDIOTYPE
- 9) WEI: CRYOGLOBULIN WITH ANTI-LOW-DENSITY LIPOPROTEIN ACTIVITY; B IDIOTYPE
- 10) GAR: CRYOGLOBULIN WITH ANTI-HGG ACTIVITY; B IDIOTYPE
- 11) PIE: AUTOCARDIOTYPIC WHICH BINDS SPECIFICALLY TO INTERMEDIATE FILAMENTS
- 12) FLO: CRYOGLOBULIN WITH ANTI-HGG ACTIVITY; B IDIOTYPE
- 13) LOP: CRYOGLOBULIN WITH ANTI-HGG ACTIVITY; B IDIOTYPE
- 14) SCA: CRYOGLOBULIN WITH ANTI-LOW-DENSITY LIPOPROTEIN ACTIVITY; B IDIOTYPE
- 15) GLO: ANTI-HUMAN GAMMA G GLOBULIN; WA IDIOTYPE; CRYOGLOBULIN WITH ANTI-HGG ACTIVITY; B IDIOTYPE
- 18) MA: COLD AGGLUTININ WITH ANTI-BLOOD GROUP I ACTIVITY (GROUP 1)
- 19) NIC: COLD AGGLUTININ WITH ANTI-BLOOD GROUP SMALL I ACTIVITY
- 20) CUR: CRYOGLOBULIN WITH ANTI-HGG ACTIVITY; B IDIOTYPE
- 22) DRE: COLD AGGLUTININ WITH ANTI-BLOOD GROUP I ACTIVITY
- 23) PER: COLD AGGLUTININ WITH ANTI-BLOOD GROUP I ACTIVITY
- 26) STE: COLD AGGLUTININ WITH ANTI-BLOOD GROUP I ACTIVITY
- 28) QJ: COLD AGGLUTININ WITH ANTI-BLOOD GROUP I ACTIVITY (ATYPICAL)
- 29) TAK: COLD AGGLUTININ WITH ANTI-BLOOD GROUP I ACTIVITY
- 35) AJ: COLD AGGLUTININ WITH ANTI-BLOOD GROUP I ACTIVITY
- 42) CLA: CRYOGLOBULIN WITH ANTI-HGG ACTIVITY; B IDIOTYPE
- 43) SHE: CRYOGLOBULIN WITH ANTI-HGG ACTIVITY; B IDIOTYPE
- 48) POM: ANTI-HUMAN GAMMA G1 GLOBULIN; PO IDIOTYPE
- 54) GOEI: ANTI-MEASLES VIRUS (WOODFOLK STRAIN); ANTI-SUBACUTE SCLEROSING PANENCEPHALITIS VIRUS (LEC STRAIN)
- 62) TEH: ANTI-HUMAN GAMMA G GLOBULIN
- 63) CRA(III): ANTI-HUMAN GAMMA G GLOBULIN
- 64) PLA: ANTI-HUMAN GAMMA G GLOBULIN
- 65) PIN: ANTI-HUMAN GAMMA G GLOBULIN
- 70) BOR: COLD AGGLUTININ WITH ANTI-BLOOD GROUP I ACTIVITY
- 71) DRI: ANTI-HUMAN GAMMA G GLOBULIN
- 72) WAL: ANTI-HUMAN GAMMA G GLOBULIN
- 73) GOL: ANTI-HUMAN GAMMA G GLOBULIN
- 74) QAG: COLD AGGLUTININ WITH ANTI-BLOOD GROUP I ACTIVITY

CLASS: HUMAN KAPPA LIGHT CHAINS SUBGROUP III

- 5) NEU: IGM-KAPPA
- 6) GOT: IGM-KAPPA
- 7) PAY: IGM-KAPPA
- 8) SON: IGM-KAPPA
- 9) WEI: IGM-KAPPA
- 10) GAR: IGM-KAPPA
- 11) PIE: IGM-KAPPA
- 12) FLO: IGM-KAPPA
- 13) LOP: IGM-KAPPA
- 14) SCA: IGM-KAPPA
- 15) GLO: IGM-KAPPA
- 20) CUR: IGM-KAPPA
- 42) CLA: IGM-KAPPA
- 43) SHE: IGM-KAPPA

REFERENCE: HUMAN KAPPA LIGHT CHAINS SUBGROUP III

- 1) TI: SUTER L. BARNIKOL H.U. WATANABE S. & HILSCHMANN N. (1969) Z. PHYSIOL. CHEM. 350, 275-278; (1972) Z. PHYSIOL. CHEM. 353, 189-208. (CHECKED BY AUTHOR)
- 2) WOL: ANDREWS D.W. & CAPRA J.D. (1981) PROC. NAT. ACAD. SCI. USA 78, 3799-3803. (CHECKED BY AUTHOR 08/25/81); ANDREWS D.W. & CAPRA J.D. (1981) BIOCHEMISTRY 20, 5816-5822.
- 3) SIE: CAPRA J.D. (1975) ADV. IMMUNOL. 20, 1-40. (CHECKED BY AUTHOR); ANDREWS D.W. & CAPRA J.D. (1981) PROC. NAT. ACAD. SCI. USA 78, 3799-3803. (CHECKED BY AUTHOR 08/25/81) WHO SUGGESTED THAT THE SEQUENCE DETERMINED IN 1975 WAS INCORRECT AND SHOULD BE DELETED; ANDREWS D.W. & CAPRA J.D. (1981) BIOCHEMISTRY 20, 5816-5822.
- 4) NG/SCL: BENTLEY D.L. (1984) NATURE 307, 77-80.
- 5) NEU: LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1983) J. IMMUNOL. 131, 1322-1325. (CHECKED BY AUTHOR 03/23/84); GONIF. F. CHEN P.P. PONS-ESTEL B. CARSON D.A. & FRANGIONE B. (1985) J. IMMUNOL. 135, 4073-4079.
- 6) GOT: LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1983) J. IMMUNOL. 131, 1322-1325. (CHECKED BY AUTHOR 03/23/84); PONS-ESTEL B. GONIF. F. SOLOMONA A. & FRANGIONE B. (1984) J. EXP. MED. 160, 859-904; GONIF. F. CHEN P.P. PONS-ESTEL B. CARSON D.A. & FRANGIONE B. (1985) J. IMMUNOL. 135, 4073-4079.
- 7) PAY: LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1983) J. IMMUNOL. 131, 1322-1325. (CHECKED BY AUTHOR 03/23/84); GONIF. F. CHEN P.P. PONS-ESTEL B. CARSON D.A. & FRANGIONE B. (1985) J. IMMUNOL. 135, 4073-4079.
- 8) SON: LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1983) J. IMMUNOL. 131, 1322-1325. (CHECKED BY AUTHOR 03/23/84); PONS-ESTEL B. GONIF. F. SOLOMONA A. & FRANGIONE B. (1984) J. EXP. MED. 160, 859-904; GONIF. F. CHEN P.P. PONS-ESTEL B. CARSON D.A. & FRANGIONE B. (1985) J. IMMUNOL. 135, 4073-4079.
- 9) WEI: LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1983) J. IMMUNOL. 131, 1322-1325. (CHECKED BY AUTHOR 03/23/84); LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1983) J. IMMUNOL. 131, 1322-1325. (CHECKED BY AUTHOR 03/23/84); PONS-ESTEL B. GONIF. F. SOLOMONA A. & FRANGIONE B. (1984) J. EXP. MED. 160, 859-904; GONIF. F. CHEN P.P. PONS-ESTEL B. CARSON D.A. & FRANGIONE B. (1985) J. IMMUNOL. 135, 4073-4079.
- 10) GAR: LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1983) J. IMMUNOL. 131, 1322-1325. (CHECKED BY AUTHOR 03/23/84); GONIF. F. CHEN P.P. PONS-ESTEL B. CARSON D.A. & FRANGIONE B. (1985) J. IMMUNOL. 135, 4073-4079.
- 11) PIE: LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1984) J. EXP. MED. 160, 893-904. (CHECKED BY AUTHOR 05/16/86)
- 12) FLO: LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1983) J. IMMUNOL. 131, 1322-1325. (CHECKED BY AUTHOR 03/23/84); GONIF. F. CHEN P.P. PONS-ESTEL B. CARSON D.A. & FRANGIONE B. (1985) J. IMMUNOL. 135, 4073-4079.
- 13) LOP: LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1983) J. IMMUNOL. 131, 1322-1325. (CHECKED BY AUTHOR 03/23/84)
- 14) SCA: LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1983) J. IMMUNOL. 131, 1322-1325. (CHECKED BY AUTHOR 03/23/84)
- 15) GLO: CAPRA J.D. (1975) ADV. IMMUNOL. 20, 1-40. (CHECKED BY AUTHOR); LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1983) J. IMMUNOL. 131, 1322-1325. (CHECKED BY AUTHOR 03/23/84); GONIF. F. CHEN P.P. PONS-ESTEL B. CARSON D.A. & FRANGIONE B. (1985) J. IMMUNOL. 135, 4073-4079.
- 16) SAL: CAPRA J.D. KEHOE J.M. WINCHESTER R.J. & KUNKEL H.G. (1971) ANN. N.Y. ACAD. SCI. 190, 371-381. (CHECKED BY AUTHOR)
- 17) WIL: CAPRA J.D. KEHOE J.M. WINCHESTER R.J. & KUNKEL H.G. (1971) ANN. N.Y. ACAD. SCI. 190, 371-381. (CHECKED BY AUTHOR)
- 18) MA: CAPRA J.D. KEHOE J.M. WILLIAMS R.C. JR. FEIZT. & KUNKEL H.G. (1972) PROC. NAT. ACAD. SCI. USA 69, 40-43. (CHECKED BY AUTHOR)
- 19) NIC: CAPRA J.D. KEHOE J.M. WILLIAMS R.C. JR. FEIZT. & KUNKEL H.G. (1972) PROC. NAT. ACAD. SCI. USA 69, 40-43. (CHECKED BY AUTHOR)
- 20) CUR: LEDFORD D.K. GONIF. PIZZOLATO M. FRANKLINE E.C. SOLOMONA A. & FRANGIONE B. (1983) J. IMMUNOL. 131, 1322-1325. (CHECKED BY AUTHOR 03/23/84); GONIF. F. CHEN P.P. PONS-ESTEL B. CARSON D.A. & FRANGIONE B. (1985) J. IMMUNOL. 135, 4073-4079.
- 21) FRA: MILSTEIN C. (1969) FEBS LETTERS 5, 301-304. (CHECKED BY AUTHOR WHO PROVIDED ADDITIONAL RESIDUES TO THOSE PUBLISHED)
- 22) DRE: GERGELY J., WANG A.C. & FUDENBERG H.H. (1973) VOX SANG. 24, 432-440. (CHECKED BY AUTHOR)
- 23) PER: GERGELY J., WANG A.C. & FUDENBERG H.H. (1973) VOX SANG. 24, 432-440. (CHECKED BY AUTHOR)
- 24) CAM: HOPFER J.E. NOYES C., HSUR H., HEINRIKSON R. & GALLAGHER W. (1979) J. IMMUNOL. 122, 2007-2010. (CHECKED BY AUTHOR 01/26/83)
- 25) STE: EDMAN M. & COOPER G. (1968) FEBS LETTERS 2, 33-35. (CHECKED BY AUTHOR)
- 26) QJ: CAPRA J.D. KEHOE J.M. WILLIAMS R.C. JR. FEIZT. & KUNKEL H.G. (1972) PROC. NAT. ACAD. SCI. USA 69, 40-43. (CHECKED BY AUTHOR)
- 27) TAK: GERGELY J., WANG A.C. & FUDENBERG H.H. (1973) VOX SANG. 24, 432-440. (CHECKED BY AUTHOR)
- 28) IARC/BL41: KLOBECK H.G. MEINDL A., COMBIATO G. SOLOMONA A. & ZACHAU H.G. (1985) NUCLEIC ACIDS RES. 13, 6499-6513.
- 29) RAD: MILSTEIN C. (1969) FEBS LETTERS 5, 301-304. (CHECKED BY AUTHOR)
- 30) DIL: DAYHOFF M.O. (1972) ATLAS OF PROTEIN SEQUENCE & STRUCTURE, D-250. SUBMITTED BY SMITHIES G., GIBSON D.M. AND FANNING E.M. (CHECKED BY AUTHOR 07/24/79)

REFERENCE: HUMAN KAPPA LIGHT CHAINS SUBGROUP III (cont'd)

- 31) CAS: NALL.H.D. & EDMAN.P. (1967) NATURE 216:262-263. (CHECKED BY AUTHOR 07/25/79)
- 32) MCE: MIDDUGH.G.R., KEHOE.J.M., PRYSTOWSKY.M.B., GERBER-JENSON.S., JENSON.J.C. & LITMAN.G.W. (1978) IMMUNOCHEM. 15:171-187. (CHECKED BY AUTHOR 10/22/80)
- 33) KAE: WANG.A.C. & FUDENBERG.H.H. (1975) IMMUNOL.COMMUN. 4:483-487. (CHECKED BY AUTHOR 09/23/77); WANG.A.C., TUNG.E., WANG.I., FUDENBERG.H.H., PICK.A.I. & FROELICHMAN.R. (1980) CANCER IMMUNOL.IMMUNOTHER. 9:81-86. (CHECKED BY AUTHOR 03/19/81)
- 34) SMJ: NALL.H.D. & EDMAN.P. (1967) NATURE 216:262-263. (CHECKED BY AUTHOR 07/25/79)
- 41) CAPRA.J.D., KEHOE.J.M., WILLIAMS.R.C., JR., FEIZ.T. & KUNKEL.H.G. (1972) PROC.NAT.ACAD.SCI.USA 69:40-43. (CHECKED BY AUTHOR)
- 60) BROGGI: HOPPER.J.E., MOVES.C., HENDERSON.E.S. & PRESSMAN.D. (1977) IMMUNOCHEM. 14:567-572. (CHECKED BY AUTHOR 01/26/83)
- 37) NIG: NALL.H.D. & EDMAN.P. (1967) NATURE 216:262-263. (CHECKED BY AUTHOR 07/25/79)
- 38) IKE: CAPRA.J.D. (1975) ADV.IMMUNOLOGY 20:1-40. (CHECKED BY AUTHOR)
- 39) TIL: PINK.I.R.L., WANG.A.C. & FUDENBERG.H.H. (1971) ANN.REV.MED. 22:145-170. (CHECKED BY AUTHOR)
- 40) AMYLOID KSA: SLETTEN.K., WESTERMARK.P., PITKANEN.P., THYRESSON.N. & OLSTAD.O.K. (1983) SCAND.J.IMMUNOL. 18:557-560. (CHECKED BY AUTHOR 04/26/84)
- 41) POL: WANG.A.C., WELLS.J.V., FUDENBERG.H.H. & GERGELY.J. (1974) IMMUNOCHEM. 11:341-345. (CHECKED BY AUTHOR)
- 42) CLA: LEDFORD.D.K., GONI.F., PIZZOLATO.M., FRANKLIN.E.C., SOLOMON.A. & FRANGIONE.B. (1983) J.IMMUNOL. 131:1322-1325. (CHECKED BY AUTHOR 03/23/84)
- 43) SHE: LEDFORD.D.K., GONI.F., PIZZOLATO.M., FRANKLIN.E.C., SOLOMON.A. & FRANGIONE.B. (1983) J.IMMUNOL. 131:1322-1325. (CHECKED BY AUTHOR 03/23/84)
- 44) JH: JEMMERSON.R., KAPLAN.B., DENTON.N.D., ANDERAS.P., ANDERSON.B. & MARGOLASH.E. (1979) BIOCHEMISTRY 18:4676-4683.
- 45) WIN: NALL.H.D. & EDMAN.P. (1967) NATURE 216:262-263. (CHECKED BY AUTHOR 07/25/79)
- 46) LEA: WANG.A.C., WELLS.J.V., FUDENBERG.H.H. & GERGELY.J. (1974) IMMUNOCHEM. 11:341-345. (CHECKED BY AUTHOR)
- 47) ARP: WANG.A.C., TUNG.E., WANG.I., FUDENBERG.H.H., PICK.A.I. & FROELICHMAN.R. (1980) CANCER IMMUNOL.IMMUNOTHER. 9:81-86. (CHECKED BY AUTHOR 03/19/81)
- 48) POM: KLAPPER.D.G. & CAPRA.J.D. (1978) ANN.IMMUNOL.(INST.PASTEUR) 127C:261-271. (CHECKED BY AUTHOR 08/07/79)
- 49) VAND: SEON.B.K., GAILANI.S., HENDERSON.E.S. & PRESSMAN.D. (1977) IMMUNOCHEM. 14:567-572. (CHECKED BY AUTHOR 08/27/80)
- 50) AMYLOID S0124: SLETTEN.K., WESTERMARK.P., PITKANEN.P., THYRESSON.N. & OLSTAD.O.K. (1983) SCAND.J.IMMUNOL. 18:557-560. (CHECKED BY AUTHOR 04/26/84)
- 51) DOY: WANG.A.C., TUNG.E., WANG.I., FUDENBERG.H.H., PICK.A.I. & FROELICHMAN.R. (1980) CANCER IMMUNOL.IMMUNOTHER. 9:81-86. (CHECKED BY AUTHOR 03/19/81)
- 52) SHM: WANG.A.C., TUNG.E., WANG.I., FUDENBERG.H.H., PICK.A.I. & FROELICHMAN.R. (1980) CANCER IMMUNOL.IMMUNOTHER. 9:81-86. (CHECKED BY AUTHOR 03/19/81)
- 53) GRA: NALL.H.D. & EDMAN.P. (1967) NATURE 216:262-263. (CHECKED BY AUTHOR 07/25/79)
- 54) GORJ: STROBERG.A.D., KARCHER.D. & LOWENTHAL.A. (1975) J.IMMUNOL. 115:157-160. (CHECKED BY AUTHOR)
- 55) LOW: WANG.A.C., TUNG.E., WANG.I., FUDENBERG.H.H., PICK.A.I. & FROELICHMAN.R. (1980) CANCER IMMUNOL.IMMUNOTHER. 9:81-86. (CHECKED BY AUTHOR 03/19/81)
- 56) VER: CHERIAU.A. & NATALI.P.G. (1978) IMMUNOCHEMISTRY 15:585-589. (CHECKED BY AUTHOR 09/13/79)
- 57) REE: PRELLIF., TUMMLO.D., SOLOMON.A. & FRANGIONE.B. (1986) J.IMMUNOL. IN PRESS.
- 58) WE: DWORSKY.E., SLETTEN.K., HARBOE.M. & WETTELAND.P. (1980) SCAND.J.IMMUNOL. 12:281-287. (CHECKED BY AUTHOR 02/28/1984)
- 59) HOW: KAPLAN.A.P. & METZGER.H. (1969) BIOCHEMISTRY 8:3944-3951. (CHECKED BY AUTHOR)
- 60) HS4: HOOD.L., GRAY.W.R., SANDERS.B.G. & DREYER.W.J. (1967) COLD SPRING HARBOR SYMP. QUANTITATIVE BIOL. 32:133-145.
- 61) HS5: HOOD.L., GRAY.W.R., SANDERS.B.G. & DREYER.W.J. (1967) COLD SPRING HARBOR SYMP. QUANTITATIVE BIOL. 32:133-145.
- 62) TEH: JOHNSTON.S.L., ABRAHAM.G.N. & WELCH.E.H. (1975) BIOCHEM.BIOPHYS.RES.COMMUN. 66:842-847. (CHECKED BY AUTHOR 10/17/77)
- 63) CRA(II): JOHNSTON.S.L., ABRAHAM.G.N. & WELCH.E.H. (1975) BIOCHEM.BIOPHYS.RES.COMMUN. 66:842-847. (CHECKED BY AUTHOR 10/17/77)
- 64) PLA: JOHNSTON.S.L., ABRAHAM.G.N. & WELCH.E.H. (1975) BIOCHEM.BIOPHYS.RES.COMMUN. 66:842-847. (CHECKED BY AUTHOR 10/17/77)
- 65) PIN: JOHNSTON.S.L., ABRAHAM.G.N. & WELCH.E.H. (1975) BIOCHEM.BIOPHYS.RES.COMMUN. 66:842-847. (CHECKED BY AUTHOR 10/17/77)
- 66) MCE: CAPRA.J.D., KEHOE.J.M., WILLIAMS.R.C., JR., FEIZ.T. & KUNKEL.H.G. (1972) PROC.NAT.ACAD.SCI.USA 69:40-43. (CHECKED BY AUTHOR)
- 67) HAC: HOOD.L. & TALMAGE.D.W. (1970) SCIENCE 166:325-334.
- 68) K. EVIS'CL: STAVNEZER.J., KEKISH.O., BATTER.D., GRENIER.J., BALAZS.I., HENDERSON.E. & ZEGERS.B.J.M. (1985) NUC.ACIDS RES. 13:3495-3514.
- 69) BER: WANG.A.C., WELLS.J.V., FUDENBERG.H.H. & GERGELY.J. (1974) IMMUNOCHEM. 11:341-345. (CHECKED BY AUTHOR)
- 70) BOR: GERGELY.J., WANG.A.C. & FUDENBERG.H.H. (1973) VOK SANG 24:432-440. (CHECKED BY AUTHOR)
- 71) DRI: CAPRA.J.D. (1975) ADV.IMMUNOLOGY 20:1-40. (CHECKED BY AUTHOR)
- 72) WAL: CAPRA.J.D. (1975) ADV.IMMUNOLOGY 20:1-40. (CHECKED BY AUTHOR)
- 73) GOL: CAPRA.J.D. (1975) ADV.IMMUNOLOGY 20:1-40. (CHECKED BY AUTHOR)
- 74) GAO: CAPRA.J.D., KEHOE.J.M., WILLIAMS.R.C., JR., FEIZ.T. & KUNKEL.H.G. (1972) PROC.NAT.ACAD.SCI.USA 69:40-43. (CHECKED BY AUTHOR)
- 75) DOB: HOOD.L. & TALMAGE.D.W. (1970) SCIENCE 166:325-334.
- 76) HS6: HOOD.L., GRAY.W.R., SANDERS.B.G. & DREYER.W.J. (1967) COLD SPRING HARBOR SYMP. QUANTITATIVE BIOL. 32:133-145.
- 77) HS12: HOOD.L., GRAY.W.R., SANDERS.B.G. & DREYER.W.J. (1967) COLD SPRING HARBOR SYMP. QUANTITATIVE BIOL. 32:133-145.
- 78) BURK: MOULINA.A. & FOUGERE.A.M. (1973) NATURE NEW BIOLOGY 246:178-178. (CHECKED BY AUTHOR)
- 79) ED: MOULINA.A. & FOUGERE.A.M. (1973) NATURE NEW BIOLOGY 246:178-178. (CHECKED BY AUTHOR)
- 80) BE: MILSTEIN.C. (1969) FEBS LETTERS 2:301-304. (CHECKED BY AUTHOR)
- 81) AMYLOID WR: WESTERMARK.P., SLETTEN.K., PITKANEN.P., NATVIG.J.B. & LINDHOLM.C.E. (1982) MOL.IMMUNOL. 19:447-450. (CHECKED BY AUTHOR 05/01/83)
- 82) VKAPPA3'CL: BENTLEY.D.L. & RABBITS.T.H. (1981) CELL 24:613-623. (CHECKED BY AUTHOR 12/07/81)

NOTES: HUMAN KAPPA LIGHT CHAINS SUBGROUP III

IDENTICAL SETS OF FRAMEWORK SEGMENTS:

- FR1: SET 1: T1[1], WOL2[1], SIE3[1], N99[CL4], NEU5[5], G0T6[8], S0N8[8], WEI[8], G1AR[10], PIE1[1], FLO1[12], LOP1[13], SCA1[4], GLO1[15], SAL1[16], WU1[17], MA1[18], NIC1[19], CUR1[20], FR4[21], DRE1[22], PER1[23], CAM1[24], (24 IDENTICAL)
- SET 2: GA1[26], TAK1[27], (2 IDENTICAL)
- SET 3: RA1[29], CL1[30], CAS1[31], WEL1[32], (4 IDENTICAL)
- SET 4: KEA1[33], SM1[34], (2 IDENTICAL HUMAN V-KAPPA-III; ALSO 1 MOUSE V-KAPPA-V: Vg CL122)
- SET 5: K1A1[35], BRO1[36], (2 IDENTICAL)
- SET 6: CL4[42], SHE1[43], (2 IDENTICAL)
- FR2: SET 1: T1[1], WOL2[1], SIE3[1], N99[CL4], NEU5[5], G0T6[8], S0N8[8], GAR1[10], PIE1[1], FLO1[12], GLO1[15], CUR1[20], (12 IDENTICAL HUMAN V-KAPPA-III; ALSO 1 MOUSE V-KAPPA-IV: Vg CL121); AND 1 MOUSE V-KAPPA-V: Vg CL122)
- FR3: SET 1: T1[1], WOL2[1], (2 IDENTICAL)
- SET 2: G0T6[8], PAY1[7], SASA[10], PIE1[1], FLO1[12], GLO1[15], CUR1[20], (7 IDENTICAL)
- FR4: SET 1: WOL2[1], PAY1[7], PIE1[1], GLO1[15], CUR1[20], REE1[57], VKAPPA3'CL[82], (7 IDENTICAL HUMAN V-KAPPA-III; ALSO 3 HUMAN V-KAPPA-IV: AU2[2], GAL1[36], CL1[11]; 2 HUMAN V-KAPPA-III: GM 607 [CL5], RPM1-6410 [CL16]; AND 1 HUMAN V-KAPPA-IV: PB17V [CL3])
- SET 2: POM1[48], (IDENTICAL TO 1 HUMAN V-KAPPA-III: HAU1[4])
- SET 3: NEU5[5], G0T6[8], GAR1[10], FLO1[12], FR4[21], IARC[BL41] CL28[1], (6 IDENTICAL HUMAN V-KAPPA-III; ALSO 3 HUMAN V-KAPPA-IV: AG17[1], DEN1[46], BIE3[1]; 2 HUMAN V-KAPPA-III: NMJ3 [FR14]; AND 1 HUMAN V-KAPPA-IV: LENA1[1])
- SET 4: S0N8[8], (IDENTICAL TO 1 HUMAN V-KAPPA-IV: VJ[CL11])
- IDENTICAL SETS OF COMPLEMENTARITY DETERMINING REGIONS:
- CDR1: SET 1: SIE3[1], KE1[38], (2 IDENTICAL)
- SET 2: N99[CL4], PAY1[7], S0S1[WEI[9], GAR1[10], PIE1[1], FLO1[12], GLO1[15], CUR1[20], DRE1[22], CAM1[24], (11 IDENTICAL)
- SET 3: TIL3[39], (IDENTICAL TO 1 MOUSE V-KAPPA-V: Vg CL122)
- CDR2: SET 1: SIE3[1], SIE3[1], NEU5[5], G0T6[8], GAR1[10], FLO1[12], GLO1[15], CUR1[20], (10 IDENTICAL)
- SET 2: POM1[48], (IDENTICAL TO 1 MOUSE V-KAPPA-IV: Vg CL121)
- CDR3: SET 1: POM1[48], (IDENTICAL TO 1 HUMAN V-KAPPA-III: LAY1[39])
- SET 2: G0T6[8], CUR1[20], (2 IDENTICAL)
- SET 3: PAY1[7], GLO1[15], (2 IDENTICAL)
- SET 4: GAR1[10], FLO1[12], (2 IDENTICAL)
- IDENTICAL SETS OF J-MINIGENES:
- SET 1: PIE1[1], VKAPPA3'CL[82], (2 IDENTICAL HUMAN V-KAPPA-III; ALSO 1 HUMAN V-KAPPA-IV: AU2[2]; 1 HUMAN V-KAPPA-III: RPM1-6410 [CL16]; AND 1 HUMAN V-KAPPA-IV: PB17V [CL3])
- SET 2: G0T6[8], (IDENTICAL TO 1 HUMAN V-KAPPA-III: AG17[1])
- SET 3: GAR1[10], FLO1[12], IARC[BL41] CL28[1], (3 IDENTICAL HUMAN V-KAPPA-III; ALSO 2 HUMAN V-KAPPA-III: DEN1[46], BIE3[1]; AND 1 HUMAN V-KAPPA-IV: FR1[4])
- SET 4: WOL2[1], CUR1[20], (2 IDENTICAL)
- SET 5: PAY1[7], GLO1[15], (2 IDENTICAL)

SPECIFIC NOTES:

- 41) NG9'CL: THE AMINO ACID SEQUENCE IS TRANSLATED FROM THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN CDNA.
- 32) MCE: IT IS A CRYOIMMUNOGLOBULIN. THE AUTHORS ORIGINALLY DESIGNATED IT AS MCE, BUT IN ORDER TO DIFFERENTIATE IT FROM ANOTHER MCE SEQUENCED BY CAPRA ET AL., IT IS DENOTED AS MCE.
- 42) CLA: THE AMINO ACID RESIDUES FOUND AT POSITION 9 WERE GLY AND ALA.
- 43) SHE: THE AMINO ACID RESIDUES FOUND AT POSITION 9 WERE GLY AND ALA.

NOTES: HUMAN KAPPA LIGHT CHAINS SUBGROUP III (cont'd)

- 44) JH: THE NAME WAS GIVEN TO US BY THE AUTHORS. IT IS NOT INCLUDED IN THE PAPER.
- 58) WE: AT POSITIONS 20,29 AND 33 OF AMINO ACID SEQUENCE WERE FOUND BOTH LEU AND ILE. IN THE SAME SEQUENCE TWO RESIDUES WERE FOUND IN POSITIONS 1,3,4,9,10,15,17,19,20,21,22 AND 29. THE SECOND RESIDUES WERE GLU,VAL,LEU,GLY,THR,PRO,GLU,ALA,THR,LEU,SER AND VAL. RESPECTIVELY. A DETERMINATION WAS NOT MADE IN THE ARTICLE AS TO WHETHER THE SEQUENCE BELONGED TO SUBGROUP I OR TO SUBGROUP III.
- 81) AMYLOID WR: AMINO ACID RESIDUES FOUND AT POSITION 54 ARE LEU AND ALA.
- 82) VKAPPA3⁺CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF CDNA FROM A MOUSE-HUMAN HYBRID CELL LINE.

[illegible]

	OCCURRENCES OF MOST COMMON AMINO ACID	VARIABILITY
F R I	0	
	15(ASP)	1
	14(ILE)	2.1
	15(VAL)	2.3
	13(MET)	
	14(THR)	2.1
	15(GLN)	1
	14(SER)	1
	15(PRO)	1
	10(ASP) - 7(ASP)	7.5 : 11.
	11(SER)	2.4
	14(LEU)	1
	14(ALA)	1
	14(VAL)	1
	11(SER)	1
	8(LEU)	2.8
	11(GLY)	1
	7(GLU) - 5(GLU)	3.1 : 6.6
	8(ARG)	1.1
	12(ALA)	1
O C R I	12(THR)	1
	8(ILE)	1
	7(ASN) - 4(+)	5.1 : 9.
	10(CYS)	
	5(LYS)	6.
	5(SER)	5.4
	6(SER)	2.3
	7(GLN) - 6(GLN)	1. : 2.3
	5(SER)	
	6(VAL)	
F R I	6(LEU)	
	5(TYR)	
	4(SER)	
	4(SER)	
	3(ASN)	3.3
	3(ASN)	
	5(LYS)	2.7
	4(ASN)	1
	4(TYR)	1
	4(LEU)	1
F R I	4(ALA)	
	4(THR)	1
	4(TYR)	1
	4(GLN)	1
	4(GLN)	1
	4(LYS)	1
	5(PRO)	1
	5(GLY)	1
	5(GLN)	1
	4(PRO)	2.5
O C R I	5(PRO)	1
	5(LYS)	1
	5(LEU)	1
	5(LEU)	1
	5(ILE)	1
	5(TYR)	1
	5(THR)	1
	4(THR)	1
	4(ASP)	2.5
	4(SER)	1
F R I	4(THR)	1
	4(PHE)	1
	4(THR)	1
	4(LEU)	1
	4(THR)	1
	4(ILE)	1
	4(SER)	1
	4(SER)	1
	4(LEU)	1
	4(GLN)	1
O C R I	4(ALA)	1
	4(GLU)	1
	4(ASP)	1
	4(VAL)	1
	4(VAL)	1
	4(ALA)	1
	4(VAL)	1
	4(TYR)	1
	4(TYR)	1
	4(CYS)	1
F R I	4(GLN)	1
	4(GLN)	1
	4(TYR)	1
	3(TYR)	2.7
	2(SER)	6.
	2(THR)	6.6
	4(PRO)	1
	95A	
	95B	
	95C	
O C R I	95D	
	95E	
	95F	
	1(+)	4.
	2(THR)	3.
	3(PHE)	1
	3(GLY)	1
	2(GLN)	3.
	3(GLY)	1
	3(THR)	1
F R I	3(LYS)	1
	2(+)	4.
	4(GLU)	1
	4(ILE)	1
	106A	
	3(LYS)	2.7
	3(ARG)	1
	1(THR)	
	108	
	109	

ANTIBODY SPECIFICITIES: HUMAN KAPPA LIGHT CHAINS SUBGROUP IV

- 3) PB17V-CL: ANTI-STREPTOCOCCUS GROUP A CARBOHYDRATE WITH SPECIFICITY FOR N-ACETYL GLUCOSAMINE
 5) R.K.: COLD AGGLUTININ WITH ANTI-PRH ACTIVITY (RBC MEMBRANE ANTIGEN ON HUMAN ERYTHROCYTES INACTIVATED BY PROTEOLYTIC ENZYMES AND NEURAMINIDASE)
 6) L.T.H.: COLD AGGLUTININ WITH ANTI-PRG ACTIVITY (RBC MEMBRANE ANTIGEN ON HUMAN, RAT AND GUINEA PIG ERYTHROCYTES INACTIVATED BY PROTEOLYTIC ENZYMES AND NEURAMINIDASE)
 7) TUR: COLD AGGLUTININ WITH ANTI-PR ACTIVITY

REFERENCE: HUMAN KAPPA LIGHT CHAINS SUBGROUP IV

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 11) DA-N: BOUVET,J.P.,LIACOPOULOS,P.,PILOT,J.,BANDAR,TUNG,E. & WANG,A.C. (1980) J.IMMUNOL.,125,213-220. (CHECKED BY AUTHOR 08/04/80); BOUVET,J.P.,LIACOPOULOS,P.,PILOT,J.,BANDAR,TUNG,E. & WANG,A.C. (1980) J.IMMUNOL.,129,1519-1524.
 12) JAM: SLETTEN,K.,HANNESTAD,K. & HARBOE,M. (1974) SCAND.J.IMMUNOL.,3,219-222. (CHECKED BY AUTHOR 12/05/77)
 13) SCH: SLETTEN,K.,HANNESTAD,K. & HARBOE,M. (1974) SCAND.J.IMMUNOL.,3,219-222. (CHECKED BY AUTHOR 12/05/77)
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NOTES: HUMAN KAPPA LIGHT CHAINS SUBGROUP IV

IDENTICAL SETS OF FRAMEWORK SEGMENTS:

- FR1: SET 1: VJ1-CL(1),VKAPPA IV GERMLINE-CL(2),PB17V-CL(3),R.K.(5); (4 IDENTICAL)
 SET 2: LEN(4),R.K.(5); (2 IDENTICAL)
 SET 3: DA(9),DA-H(10); (2 IDENTICAL)
 FR2: SET 1: VJ1-CL(1),VKAPPA IV GERMLINE-CL(2),PB17V-CL(3),LEN(4); (4 IDENTICAL HUMAN V-KAPPA-IV; ALSO 2 HUMAN V-KAPPA-I: V19B(1)(88); V19B(1)(89); 1 MOUSE V-KAPPA-I: MOPC-603(47); 30 MOUSE V-KAPPA-II: MPC11(1)(6); TEPC11(1)(7); PC374(1)(N2B)(9); TEPC1(24)(9); MOPC32(1)(12); PC7043(N2B)(13); PC7183(N2B)(14); PC6308(N2B)(15); PC6584(N2B)(17); PC7940(N2B)(18); PC7173(N2B)(19); PC245R(N2B)(20); PC4035(N2B)(21); PC7270(N2B)(23); K36-15(26); Z44(28); V-2151(26B)(30); V-2129(26B)(31); PC7461(N2B)(33); PC2950(N2B)(34); 97 C.A.B.Y(35); 10 A.T.H(39); H36-5(48); 40 C.A.T.H(52); MOPC63(54); ABPC22(55); PC3245(N2B)(56); PC4050(N2B)(57); V-21518(58); B1(58); 1(59); 1(59); K49-50(145); 3547(47); K4820(57); K30-26(781); 311(65); 4422(66); 17D5(1)(68); 4150(1)(1); 4363(65); 1201(5); K-25(1121)
 FR3: SET 1: VJ1-CL(1),VKAPPA IV GERMLINE-CL(2),PB17V-CL(3),LEN(4); (4 IDENTICAL)
 FR4: SET 1: PB17V-CL(3); (IDENTICAL TO 3 HUMAN V-KAPPA-I: AU1(2),GAL(1)(36); CL-1(110); 2 HUMAN V-KAPPA-II: GM 607(1)(51); RPM1-84(1)(16); AND 7 HUMAN V-KAPPA-III: WOL2(1),RAY(1),PIE(1)(1); GLQ(1)(16); CUR(20); REE(57); VKAPPA(1)(80); SET 2: LEN(4); (IDENTICAL TO 3 HUMAN V-KAPPA-I: AG17(DEN(46); B1(63); 2 HUMAN V-KAPPA-II: NIM(3); FR(14); AND 6 HUMAN V-KAPPA-III: NEU(5); GOT(16); SAR(10); FLO(12); FRA(21); IACR(4); CL(28); SET 3: VJ1-CL(1); (IDENTICAL TO 1 HUMAN V-KAPPA-III: SON(8))

IDENTICAL SETS OF COMPLEMENTARITY DETERMINING REGIONS:

- CDR1: SET 1: VJ1-CL(1),VKAPPA IV GERMLINE-CL(2); (2 IDENTICAL)
 CDR2: SET 1: VJ1-CL(1),VKAPPA IV GERMLINE-CL(2),PB17V-CL(3),LEN(4); (4 IDENTICAL HUMAN V-KAPPA-IV; ALSO 1 MOUSE V-KAPPA-IV: KPN16(1)(70))
 CDR3:

IDENTICAL SETS OF J-MINIGENES:

- SET 1: PB17V-CL(3); (IDENTICAL TO 1 HUMAN V-KAPPA-I: AU1(2); 1 HUMAN V-KAPPA-II: RPM1-84(1)(16); AND 2 HUMAN V-KAPPA-III: PIE(1)(1); VKAPPA(1)(80))

+ THE FOLLOWING WERE EQUALLY AND MOST FREQUENTLY OCCURRING:

AT POSITION	RESIDUES
22	(SER,ASP,ASN)
86	(TRP,TYR)
104	(LEU,VAL)

[illegible]

HUMAN LAMBDA LIGHT CHAINS SUBGROUP I (cont'd)

	# OF FUL #	# OF SEQUENCES	# OF AMINO ACIDS	OCCURRENCES OF MOST COMMON AMINO ACID	VARIABILITY
0					
1		20	2	18(PCA)	2.1
2		20	2	20(SER)	1.
3		21	2	20(VAL)	2.1
4		21	1	21(LEU)	1.
5		22	1	22(THR)	1.
6		21	1	21(GLN) - 20(GLN)	1. : 2.1
7		21	1	21(PRO)	1.
8		21	1	21(SER)	1.
9					
10		21	3	11(ALA)	5.7
11		22	1	22(SER)	1.
12		22	2	16(GLY)	2.8
13		22	2	11(THR)	6.
14		21	2	20(PRO)	2.1
15		21	2	21(VAL)	1.
16		21	2	20(GLN) - 19(GLN)	2.1 : 2.2
17		21	2	19(VAL)	2.1
18		20	2	16(THR)	5.
19		20	2	18(LEU)	2.1
20		19	2	18(SER)	2.1
21		19	1	19(CYS)	1.
22					
23	CYS	18	3	15(SER)	3.8
24	SER	18	1	18(GLY)	1.
25	GLY	18	1	13(SER)	3.9
26	ASN	17	3	12(SER)	6.7
27	ASN	16	5		
27A	---				
27B	---				
27C	---				
27D	SER	15	3	12(SER)	
27E		15	3	12(ASN)	
27F		4	3	2(LEU)	
28		15	5	10(LEU)	7.5
29		14	3	12(GLY)	3.5
30		14	7	4(SER)	23.
31		14	4	11(ASN)	5.1
32		14	6	5(TYR)	17.
33		14	1	14(VAL)	1.
34		14	7	4(LEU) - 3	25.
35		14	1	14(THR)	1.
36		14	2	13(TYR)	2.2
37		14	3	12(GLN)	3.5
38		14	3	9(GLN)	4.7
39		14	4	9(LEU)	6.2
40		14	1	14(PRO)	1.
41		14	1	14(GLY)	1.
42		14	3	12(THR)	3.5
43		14	2	13(ALA)	2.2
44		14	1	14(PRO)	1.
45		14	2	13(LYS)	2.2
46		14	1	14(LEU)	2.2
47		14	1	13(LEU)	2.2
48		14	2	13(LEU)	2.2
49		14	2	12(TYR)	2.3
50		14	6	4(SER)	26.
51		14	3	8(ASN)	5.3
52		14	3	8(ASN)	5.3
53		14	5	6(GLN)	12.
54		14	3	12(ARG)	3.5
55		12	3	10(PRO)	3.6
56		12	1	12(SER)	1.
57		12	1	12(GLY)	1.
58		12	2	9(VAL)	2.7
59		12	2	10(PRO)	2.4
60		12	2	11(ASP)	1.2
61		13	1	13(ARG)	2.3
62		14	2	12(PHE)	1.
63		14	1	14(SER)	1.
64		14	3	9(GLY)	4.7
65		14	1	14(SER)	1.
66		14	1	14(LYS)	1.
67		14	1	14(SER)	1.
68		14	1	14(GLY)	1.
69		14	3	12(THR)	3.5
70		14	1	14(SER)	1.
71		14	1	14(ALA)	1.
72		14	2	9(SER)	3.1
73		14	1	14(LEU)	1.
74		14	2	11(ALA)	2.5
75		14	1	14(LEU)	1.
76		14	2	2(SER)	3.1
77		14	1	14(GLY)	1.
78		14	1	14(LEU)	1.
79		14	4	9(GLN)	6.2
80		14	4	10(GLU)	7.
81		14	2	13(ASP)	2.8
82		14	2	14(GLU)	2.2
83		14	1	14(LEU)	1.
84		14	3	11(ALA)	3.8
85		14	3	12(ASP)	3.5
86		14	1	14(TYR)	1.
87		14	3	11(TYR)	1.
88		14	1	14(CYS)	3.8
89		14	3	10(ALA)	4.2
90		14	3	7(THR)	6.
91		14	2	12(TRP)	2.3
92		14	2	12(ASP)	2.3
93		14	5	8(ASP)	8.8
94		14	2	12(SER)	2.3
95		14	2	13(LEU)	2.2
95A		11	3	8(ASP)	
95B		11	4	9(GLY)	
95C					
95D					
95E					
95F		14	7	6(PRO)	18.
96		14	3	12(VAL)	3.5
97		14	1	14(PHE)	1.
98		14	1	14(GLY)	1.
99		14	2	13(GLY)	2.2
100		14	1	14(GLY)	1.
101		14	1	14(THR)	1.
102		14	1	10(LYS)	7.
103		14	5	7	4.
104		14	1	14(THR)	1.
105		14	1	14(VAL)	1.
106		14	3	12(LEU)	1.
106A					
107		14	3	11(GLY)	3.8
108		12	1	12(GLN)	1.
109		12	1	12(PRO)	1.

ANTIBODY SPECIFICITIES: HUMAN LAMBDA LIGHT CHAINS SUBGROUP I

- 1) NEWM: ANTI-[3]-HYDROXY-3,7,11,15-TETRAMETHYL HEXADECYL 2-METHYL 1,4 NAPHTHOQUINONE(VIT.K10H)
 16) KOH: ANTI-HUMAN GAMMA G GLOBULIN

REFERENCE: HUMAN LAMBDA LIGHT CHAINS SUBGROUP I

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NOTES: HUMAN LAMBDA LIGHT CHAINS SUBGROUP I

IDENTICAL SETS OF FRAMEWORK SEGMENTS:

- FR1: SET 1: WAH[71]NIG-77[8],VOR[9],RHE[10],LOC[11],OKA[12]. (6 IDENTICAL)
 FR2: SET 1: NEWM[1],ANYLOID EPS[13]. (2 IDENTICAL)
 SET 2: HA[2],NIG-64[4]. (2 IDENTICAL)
 SET 3: NIG-77[8],LOC[11]. (2 IDENTICAL)
 FR3: SET 1: NIG-64[4],BL2 'CL[6]. (2 IDENTICAL)
 FR4: SET 1: NEWM[1]. (IDENTICAL TO 1 HUMAN V-LAMBDA-II: WH[3] AND 1 HUMAN V-LAMBDA-V: BO[1])
 SET 2: NEWS[VOR[9],COX[15]. (3 IDENTICAL HUMAN V-LAMBDA-I; ALSO 1 HUMAN V-LAMBDA-V: AMYLOID-ARI[1]; AND 6 MOUSE V-LAMBDA: MOPC3[15],25],TEPC3[5],26],MAA-13[27],5-7[28],MOPC3[15-26],CL[30],MOPC3[15-37],CL[32])
 SET 3: BL2 'CL[6],RHE[10],OKA[12],NIG-51[19]. (4 IDENTICAL HUMAN V-LAMBDA-I; ALSO 5 HUMAN V-LAMBDA-II: MIB[2],LES[49],B[10],I[4],VIL[17],WIN[21]; 4 HUMAN V-LAMBDA-II: HIL[1],CAP[4],BAU[12],DEL[14]; 1 HUMAN V-LAMBDA-IV: SH[1]; 3 HUMAN V-LAMBDA-V: SUT[12],THO[4],LVB[CL[5]. AND 24 MOUSE V-LAMBDA: MOPC104[1],J558[2],XS104[3],HOPC[14],J698[5],H206[18],W3159[7],Y5431[8],Y5489[9],Y6350[10],Y5468[11],MOPC51[11],S178[13],Y5444[14],Y5066[15],S1761[16],H2029[17],RJC20[18],IG
 YOL[LAMBDA-CL[19],S43[CL[21],S2H5[CL[38],S2E9[CL[39],S1F12[CL[40],IG 25LAMBDA CL[41])
 SET 4: LOC[11]. (IDENTICAL TO 1 HUMAN V-LAMBDA-V: MCB[3])

IDENTICAL SETS OF COMPLEMENTARITY DETERMINING REGIONS:

- CDR1: SET 1: NIG-64[4],BL2 'CL[6]. (2 IDENTICAL)
 CDR2: SET 1: VOR[9],NIG-51[19]. (2 IDENTICAL)
 CDR3: SET 1: VOR[9],NIG-51[19]. (2 IDENTICAL)

IDENTICAL SETS OF J-MINGENES:

- SET 1: NEWS[VOR[9],COX[15]. (2 IDENTICAL TO 1 HUMAN V-LAMBDA-V: AMYLOID-ARI[1])
 SET 2: BL2 'CL[6]. (IDENTICAL TO 2 HUMAN V-LAMBDA-V: SUT[12],THO[4]; AND 24 MOUSE V-LAMBDA: MOPC104[1],J558[2],XS104[3],HOPC[14],J698[5],H206[18],W3159[7],Y5431[8],Y5489[9],Y6350[10],Y5468[11],MOPC51[11],S178[13],Y5444[14],Y5066[15],S1761[16],H2029[17],RJC20[18],IG
 SET 3: VOR[9],COX[15]. (2 IDENTICAL)
 SET 4: HA[2],NIG-51[19]. (2 IDENTICAL)

SPECIFIC NOTES:

- 24) FUL: SOX AND HOOD HAVE REPORTED FOUR HUMAN V KAPPA AND ONE V LAMBDA CHAINS WITH ASN-SER/THR TO CONTAIN CARBOHYDRATE.

- THE FOLLOWING WERE EQUALLY AND MOST FREQUENTLY OCCURRING:

AT POSITION	RESIDUES
34	(SER,ASN)
104	(LEU,VAL)

[illegible]

HUMAN LAMBDA LIGHT CHAINS SUBGROUP II (cont'd)

	25 WAL	26 ALA	# OF SEQUENCES	# OF AMINO ACIDS	OCCURRENCES OF MOST COMMON AMINO ACID	VARIABILITY
0
1	PCA	gln	26	3	24(PCA)	3.3
2	SER	thr	26	2	25(SER)	2.1
3	val	val	26	2	23(ALA)	3.4
4	LEU	val	26	2	25(LEU)	2.1
5	THR	THR	26	2	23(THR)	3.4
6	GLN	GLN	26	1	26(GLN) 25(GLN)	1. 2.1
7	PRO	glu	26	3	24(PRO)	3.3
8	PRO	PRO	26	3	18(ALA)	3.3
9	SER	SER	26	2	25(SER)	2.1
10
11	ala	leu	26	3	23(VAL)	3.4
12	SER	thr	26	2	25(SER)	2.1
13	GLY	val	26	4	23(GLY)	4.5
14	thr	SER	26	2	25(SER)	2.1
15	PRO	PRO	26	2	25(PRO)	2.1
16	GLY	GLY	26	1	26(GLY)	1
17	GLN	glu	26	3	23(GLN)	4.5
18	arg	thr	26	3	23(SER)	3.4
19	...	val	25	3	18(LEU)	4.2
20	THR	...	25	1	25(THR)	...
21	leu	...	19	1	17(LEU)	3.4
22	thr	...	18	2	17(SER)	...
23	CYS	...	18	1	18(CYS)	...
24	ALA	...	15	4	9(THR)	6.7
25	SER	...	15	2	14(GLY)	2.1
26	SER	...	15	5	10(THR)	7.5
27	THR	...	15	5	7(SER)	11.
27A
27B
27C
27D	GLY	...	15	4	12(SER)	...
27E	ALA	...	15	4	11(ASP)	...
27F	VAL	...	15	2	14(VAL)	...
28	THR	...	15	2	10(GLY)	7.5
29	SER	...	14	5	6(GLY)	12.
30	GLY	...	14	6	8(TYR)	9.3
31	TYR	...	14	5	8(ASN) 7(ASN)	12. 14.
32	TYR	...	14	5	5(TYR)	14.
33	PRO	...	13	3	11(VAL)	3.5
34	ASN	...	13	2	12(SER)	2.2
35	TRP	...	14	1	14(TRP)	...
36	PHE	...	14	2	10(TYR)	2.8
37	GLN	...	14	2	14(GLN) 13(GLN)	2.2
38	14	2	13(GLN) 11(GLN)	2.2 3.5
39	LYS	...	14	5	10(HIS)	7.
40	PRO	...	14	1	14(PRO)	...
41	GLY	...	14	1	13(GLY)	2.2
42	GLN	...	14	4	11(LYS)	5.1
43	ALA	...	14	4	14(LEU)	2.2
44	PRO	...	14	1	14(PRO)	...
45	ARG	...	14	2	13(LYS)	2.2
46	ALA	...	14	3	12(LEU)	3.5
47	LEU	...	14	1	2()	8.4
48	ILE	...	14	1	14(ILE)	...
49	TYR	...	14	3	9(TYR)	4.7
50	SER	...	14	5	7(ASP)	10.
51	THR	...	14	4	11(VAL)	5.1
52	SER	...	14	5	2(SER)	14.
53	ASN	...	14	6	4()	21.
54	LYS	...	14	2	13(ARG)	2.2
55	PRO	...	14	2	13(PRO)	2.2
56	SER	...	14	1	14(SER)	...
57	TRP	...	14	2	13(GLY)	2.2
58	THR	...	14	3	10(VAL)	4.2
59	PRO	...	14	2	7(+)	4.
60	ALA	...	14	7	2(ASP)	20.
61	ARG	...	14	1	14(ARG)	...
62	SER	...	15	1	14(PHE)	2.1
63	SER	...	15	1	15(SER)	...
64	GLY	...	15	1	15(GLY)	...
65	SER	...	15	1	15(SER)	...
66	LEU	...	15	3	13(LYS)	3.5
67	LEU	...	14	2	13(SER)	2.2
68	GLY	...	14	2	12(GLY)	2.3
69	LYS	...	14	4	10(ASN) 9(ASN)	5.6 6.2
70	PHE	...	14	3	12(THR)	3.5
71	ALA	...	14	1	14(ALA)	...
72	ALA	...	14	1	13(SER)	2.2
73	LEU	...	14	1	14(LEU)	...
74	THR	...	14	1	14(THR)	...
75	LEU	...	14	2	13(LEU)	2.2
76	SER	...	14	1	14(SER)	...
77	GLY	...	14	1	14(GLY)	...
78	VAL	...	14	2	13(LEU)	2.2
79	GLN	...	14	3	12(GLN)	3.5
80	PRO	...	14	3	10(ALA)	4.2
81	GLU	...	14	2	11(GLU)	3.8
82	ASP	...	14	2	13(ASP)	...
83	GLU	...	14	1	14(GLU)	...
84	ALA	...	14	1	14(ALA)	...
85	GLU	...	14	3	11(ASP) 10(ASP)	3.8 5.6
86	TYR	...	14	2	14(TYR)	...
87	TYR	...	14	2	12(TYR)	2.3
88	CYS	...	14	1	14(CYS)	...
89	LEU	...	14	4	8(SER)	7.
90	LEU	...	14	2	13(SER)	2.2
91	TYR	...	14	2	12(TYR)	...
92	TYR	...	14	7	5(ALA)	20.
93	GLY	...	14	4	7(GLY)	8.
94	GLY	...	14	5	5(SER)	14. 17.
95	ALA	...	13	7	3(+)	30.
95A	11	3	2(+)	...
95B	2	2	1(+)	...
95C
95D
95E
95F
96	VAL	...	13	8	5(VAL)	21.
97	VAL	...	16	3	10(VAL)	4.8
98	PHE	...	16	1	16(PHE)	...
99	GLY	...	18	1	18(GLY)	...
100	SER	...	18	4	10(GLY)	7.2
101	GLY	...	18	1	18(GLY)	...
102	THR	...	18	1	18(THR)	...
103	LYS	...	18	5	13(LYS)	6.9
104	VAL	...	15	2	9(LEU)	3.3
105	THR	...	15	3	13(THR)	3.5
106	THR	...	13	1	13(VAL)	...
106A	13	1	13(LEU)	...
107	13	1	8(GLY)	4.9
108	10	1	10(GLN) 9(GLN)	1. 2.2
109	10	2	10(PRO)	...

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NOTES: HUMAN LAMBDA LIGHT CHAINS SUBGROUP II

IDENTICAL SETS OF FRAMEWORK SEGMENTS:

- FR1: SET 1: NIG-84(1),MES(2),WH(3),NEI(4),KAR(5),RIM(6),SLA(7); (7 IDENTICAL)
SET 2: TRO(14),BOH(15); (2 IDENTICAL)
- FR2: SET 1: WH(3),BOH(15),NIG-58(16),BURI(22); (4 IDENTICAL)
- FR3:
- FR4: SET 1: WH(3); (IDENTICAL TO 1 HUMAN V-LAMBDA-I: NEWMI(1); AND 1 HUMAN V-LAMBDA-V: BO(11))
SET 2: MES(2),ES492(8),TRO(14),VIL(17),WIN(21); (5 IDENTICAL HUMAN V-LAMBDA-II; ALSO 4 HUMAN V-LAMBDA-I: SL2, CL(8),RHE(10), OKA(12),NIG-51(19); & HUMAN V-LAMBDA-III: HIL(1),CAP(4),BAU(12),DELI(14); 1 HUMAN V-LAMBDA-IV: SH(1); 3 HUMAN V-LAMBDA-V: SUT(1),THO(4),BV,CL(5); AND 24 MOUSE V-LAMBDA: MOPC(104)(1),J558(2),XS(104)(3),HOPC(14),J698(5),H206(16), W3159(7),Y54(18),Y548(9),Y5830(10),Y566(11),MOPC(11)(1),J2(5),Y544(14),Y566(15),S178(16),H202(17), PR(20)(18); 1 HUMAN V-LAMBDA-CL(19),S43,CL(21),S2H5,CL(38),S2E9,CL(39),S1F(12),CL(40),IG 25,LAMBDA,CL(41))
SET 3: NIG-84(1); (IDENTICAL TO 1 HUMAN V-LAMBDA-III: GARI(1))

IDENTICAL SETS OF COMPLEMENTARITY DETERMINING REGIONS:

- CDR1: SET 1: MES(2),VIL(17); (2 IDENTICAL HUMAN V-LAMBDA-II; ALSO 1 HUMAN V-LAMBDA-V: MCG(3))
- CDR2: SET 1: NIG-84(1),TOG(10); (2 IDENTICAL)
- CDR3:

IDENTICAL SETS OF J-MINIGENES:

- SET 1: MES(2),TRO(14); (2 IDENTICAL HUMAN V-LAMBDA-II; ALSO 1 HUMAN V-LAMBDA-III: BAU(12))
- SET 2: ES492(8),VIL(17); (2 IDENTICAL HUMAN V-LAMBDA-II; ALSO 1 HUMAN V-LAMBDA-III: DEL(14))

SPECIFIC NOTES:

- 1) SM: IT HAS O-LINKED CARBOHYDRATE ATTACHED TO SER AT POSITION 22 AND N-LINKED CARBOHYDRATE ATTACHED TO ASX AT POSITION 25.

+ THE FOLLOWING WERE EQUALLY AND MOST FREQUENTLY OCCURRING:

AT POSITION	RESIDUES
47	(LE,MET)
53	(LYS,ASN)
59	(PRO,SER)
95	(SER,ASN)
95A	(THR,SER)
95B	(LEU,ARG)

[illegible]

HUMAN LAMBDA LIGHT CHAINS SUBGROUP III (cont'd)

	23	24	25	26	27	28	# OF	# OF	OCCURRENCES	VARIABILITY
	SG	GIM	111	119	VIN	MIL	SEQUENCES	AMINO ACIDS	OF MOST COMMON AMINO ACID	
F R I	0	---	---	---	---		12	3	10(SER)	3.6
	1	TYR	TYR	TYR	GLX	TYR	27	2	26(TYR)	2.1
	2	LEU	LEU	LEU	LEU		26	6 : 7	13(GLU) : 11(GLU)	12. : 17.
	3	LEU	LEU	LEU	LEU		26	2	25(LEU)	3.2
	4	THR	THR	THR	THR		26	3	23(THR)	3.4
	5	GLN	GLX	GLN	GLX		26	1 : 2	26(GLN) : 23(GLN)	1. : 2.4
	6	PRO	PRO	PRO	PRO		25	28	23(PRO)	2.2
	8	PRO	PRO	PRO	PRO		26	1	26(PRO)	1.
	8	SER	SER				24	1	24(SER)	1.2
	10	---	---	---	---		24	3	20(VAL)	3.6
	11	TYR	VAL	VAL			24	1	24(SER)	1.
	12	SER	SER				24	2	23(VAL)	3.7
	13	VAL	VAL				22	3	18(SER)	2.1
	15						22	2	21(PRO)	1.1
	16						21	1	21(GLY)	1.
	17					met	21	1 : 2	20(GLN) : 17(GLN)	1. : 2.4
	18						20	2	19(THR)	2.1
	19						20	2	19(ALA)	14.
	20						18	6	8(ARG)	1.
	21					ILE	19	1	19(LEU)	1.
	22					THR	17	1	17(THR)	1.
	23					CYS	17	1	17(CYS)	1.
	C D R	24				GLY		17	3 : 4	13(SER)
25					GLY		18	2	15(GLY)	2.1
26					ASP		17	3	14(ASP) : 12(ASP)	3.6 : 4.3
27					GLU		15	7	5(ALA)	21.
27A										
27B										
27C										
27D										
27E										
27F										
28							16	2	13(LEU)	2.5
29							13	5 : 6	5(GLY)	13. : 16.
30							13	6 : 7	5(GLU) : 3(+)	10. : 35.
31							14	4	8(TYR)	17.
32							13	2	9(VAL)	6.5
33							13	4	8(TYR)	2.9
34							11	1	4(TYR)	11.
F R 2		35					13	1	13(THR)	1.
		36					11	1 : 2	11(GLN) : 10(GLN)	1. : 2.2
		37					11	3	9(GLN)	3.7
		38					11	2	7(LYS)	2.1
		39					10	2	9(PRO)	2.2
		40					10	1	10(GLY)	2.3
	41					9	2 : 3	8(GLN) : 7(GLN)	2.3 : 3.9	
	42					9	2	5(ALA)	3.6	
	43					10	1	10(PRO)	1.	
	44					10	3	7(VAL)	4.3	
	45					10	3	8(LEU)	4.5	
	46					10	1	10(VAL)	1.	
	47					10	2	8(LEU)	2.5	
	48					10	2	9(TYR)	2.2	
	49					10	2	10(PRO)	1.	
	50					10	5 : 6	4(GLU) : 3(GLU)	13. : 20.	
	51					11	4	4(SER)	14.	
	52					11	4	4(LYS)	11.	
	53					11	1	11(ARG)	1.	
	54					10	2	10(PRO)	2.2	
	55					10	2	9(SER)	2.2	
	56					11	3	8(GLY)	4.1	
	57					10	2	9(LEU)	2.2	
58					10	1	10(PRO)	1.		
59					11	3	9(GLU) : 8(GLU)	3.7 : 4.1		
60					11	1	11(ARG)	1.		
61					11	1	11(PHE)	1.		
62					11	1	10(SER)	1.		
63					10	2	9(GLY)	2.2		
64					10	2	9(SER)	2.2		
65					10	4	4(ASN)	10.		
66					10	1	10(SER)	1.		
67					10	1	10(GLY)	1.		
68					10	3	5(THR)	6.		
69					10	3	8(THR)	3.8		
70					10	2	8(ALA)	2.5		
71					10	3	8(THR)	3.8		
72					10	1	10(LEU)	1.		
73					10	1	10(THR)	1.		
74					10	2	9(SER)	2.2		
75					10	2	8(GLY)	2.2		
76					10	2	5(VAL)	2.5		
77					10	2	7(GLN)	2.9		
78					10	5	3(+)	17.		
79					10	1 : 2	10(ASP) : 9(ASP)	1. : 2.5		
80					10	1 : 2	10(GLU) : 9(GLU)	1. : 2.2		
81					10	1	10(ALA)	1.		
82					10	1 : 2	10(ASP) : 8(ASP)	1. : 2.5		
83					10	1	10(TYR)	1.		
84					10	2	8(TYR)	2.5		
85					10	1	10(CYS)	1.		
86					10	3	7(GLN) : 5(GLN)	4.3 : 6.		
87					10	4	4(ALA)	10.		
88					10	3	8(ASP) : 7(ASP)	3.8 : 4.3		
89					10	5	4(SER)	13.		
90					10	6	3(THR)	30.		
91					9	6	1(+)	18.		
92					2	2				
93										
94										
95A										
95B										
95C										
95D										
95E										
95F										
96						9	5	5(VAL)	9.	
97						10	3	8(VAL)	5.	
98						10	1	10(PHE)	1.	
99						11	1	11(GLY)	1.	
100						11	3	9(GLY)	3.7	
101						11	1	11(GLY)	1.	
102						11	1	11(THR)	1.	
103						11	4	8(LYS)	5.5	
104						10	2	9(LEU)	2.2	
105						10	2	9(THR)	2.2	
106						10	1	10(VAL)	1.	
107						108A	1	10(LEU)	1.	
108						8	2	6(GLY)	2.7	
109						7	1	7(GLN)	1.	
110						7	1	7(PRO)	1.	

ANTIBODY SPECIFICITIES: HUMAN LAMBDA LIGHT CHAINS SUBGROUP III

7) GAR: ANTI-RIBOFOLAVIN

REFERENCE: HUMAN LAMBDA LIGHT CHAINS SUBGROUP III

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NOTES: HUMAN LAMBDA LIGHT CHAINS SUBGROUP III

IDENTICAL SETS OF FRAMEWORK SEGMENTS:

- FR1: SET 1: HIL1[1], YO1[2], PS1[3], CAP1[4], (4 IDENTICAL)
 SET 2: LOY 1[5], LOY 6[6], (2 IDENTICAL)
- FR2:
- FR4: SET 1: HIL1[1], CAP1[4], BAU1[12], DEL1[14], (4 IDENTICAL HUMAN V-LAMBDA-III; ALSO 4 HUMAN V-LAMBDA-I; BL2[7], CL[6], RHE[10], OKA[12], NIG-511[9]; 5 HUMAN V-LAMBDA-II; MES2[2], ES492[8], TRO1[14], VIL1[17], WIN2[1]; 1 HUMAN V-LAMBDA-IV; SH1[1]; 3 HUMAN V-LAMBDA-VI; SUT1[2], THO1[4], LEV1[15], AND 24 MOUSE V-LAMBDA: MOPC10E1[1], J558[2], X5104[3], HOPC1[4], J698[5], H2061[6], V2159[7], Y3431[8], Y3485[9], Y3630[10], Y3669[11], MOPC511[12], S1781[13], Y5444[14], Y5606[15], S1761[16], H2020[17], R3C201[18], IG 30J-LAMBDA-CL1[19], S43 CL1[21], S1H5 CL1[3], S9E CL1[30], S1F1 CL1[40], IG 25-LAMBDA-CL1[41])
 SET 2: GAR1[7], (IDENTICAL TO 1 HUMAN V-LAMBDA-III; NIG-84[1])
 SET 3: KERN1[10], (IDENTICAL TO 1 HUMAN V-LAMBDA-VI; NIG-48[1])

IDENTICAL SETS OF J-MINIGENES:

- SET 1: BAU1[2], (IDENTICAL TO 2 HUMAN V-LAMBDA-II; MES2[2], TRO1[14])
 SET 2: DEL1[14], (IDENTICAL TO 2 HUMAN V-LAMBDA-II; ES492[8], VIL1[17])

SPECIFIC NOTES:

- 18) MOT: THERE ARE TWO RESIDUES IN FRONT OF POSITION 1. THEY ARE VAL AND THR.

+ THE FOLLOWING WERE EQUALLY AND MOST FREQUENTLY OCCURRING:

AT POSITION	RESIDUES
30	(ASP, ASN, GLU)
81	(MET, GLU)
94	(ILE, ARG, SER, GLY)
95A	(TYR, ALA, GLY, ASP)
95B	(HIS, GLU)

HUMAN LAMBDA LIGHT CHAINS SUBGROUP IV

	INVARIENT RESIDUES	1 2 3 4 5					# OF SEQUENCES	# OF AMINO ACIDS	OCCURRENCES OF MOST COMMON AMINO ACID	VARIABILITY
		SH	NEV	USH	PFA	FRA				
0		---	---	---	---	---	4	1	4(SER)	1.
1	SER	SER	SER	SER	SER	ala	5	2	4(GLU)	2.5
2	LEU	GLU	LEU	LEU	LEU	---	5	1	5(LEU)	1.
3		---	---	---	---	---	5	1	4(THR)	2.5
4	GLN	THR	THR	THR	THR	val	5	2	5(GLN)	3.3
5		GLN	GLN	GLN	GLN	GLN	5	3	3(PRO)	2.5
6		ASP	pro	pro	pro	pro	5	2	4(PRO)	2.5
7		PRO	PRO	PRO	PRO	ala	5	2	3(SER)	3.3
8		ALA	ser	ser	ser	ser	5	1	---	---
9		---	---	---	---	---	5	1	5(VAL)	1.
10	VAL	SER	SER	SER	SER	gla	5	2	4(SER)	2.5
11		VAL	VAL	VAL	VAL	div	5	2	4(VAL)	2.5
12		ALA	ALA	ser	ser	ALA	5	3	3(ALA)	3.3
13		---	---	---	---	---	5	1	3(PRO)	3.3
14		LEU	LEU	pro	pro	pro	5	2	5(GLY)	1.
15	GLY	GLY	GLY	GLY	GLY	GLY	5	2	5(GLN)	3.3
16		GLN	GLX	OLN	GLN	GLX	5	3	2(THR)	2.5
17		THR	THR	THR	THR	ser	5	3	2(+)	7.5
18		VAL	VAL	ala	ala	ile	5	4	3(ILE)	10.
19		ARG	ARG	ser	val	ala	5	1	4(THR)	2.5
20	ILE	ILE	ILE	ILE	ILE	gla	5	2	4(CYS)	1.
21		THR	THR	THR	THR	THR	5	1	2(SER)	6.
22	CYS	CYS	CYS	CYS	CYS	CYS	4	3	4(GLY)	1.
23		GLN	SER	SER	ILE	---	4	2	3(ASP)	2.7
24		GLY	GLY	GLY	GLY	GLY	4	1	2(+)	4.
25		ASP	ASP	ASP	ILE	---	4	2	---	---
26		SER	LYS	LYS	SER	---	4	2	---	---
27A		---	---	---	---	---	---	---	---	---
27B		---	---	---	---	---	---	---	---	---
27C		---	---	---	---	---	---	---	---	---
27D		---	---	---	---	---	---	---	---	---
27E		---	---	---	---	---	---	---	---	---
27F		---	---	---	---	---	---	---	---	---
28	LEU	LEU	LEU	LEU	ILE	---	4	2	1(ASN) : 1(ASP)	2.7
29		GLY	GLY	GLY	GLY	---	4	2	3(LEU)	1.
30	ASP	ASP	GLN	ALA	---	---	4	2	3(GLY)	18.
31	TYR	TYR	ASN	ALA	TYR	---	4	3	1(TYR)	6.
32	ASP	TYR	ASN	ALA	---	---	3	2	2(ASP)	3.
33	ALA	ALA	TYR	TYR	---	---	3	2	2(ALA)	9.
34	TRP	TRP	TRP	TRP	ILE	---	3	3	1(ILE)	1.
35	TYR	TYR	TYR	TYR	---	---	3	1	3(TYR)	1.
36	GLN	GLN	GLN	GLN	---	---	2	1	3(GLN)	1.
37	LYS	LYS	LYS	LYS	---	---	2	1	2(GLN)	1.
38		PRO	PRO	PRO	---	---	1	1	2(LYS)	1.
39		GLN	GLN	GLN	---	---	1	1	1(PRO)	---
40		ALA	ALA	ALA	---	---	1	1	1(GLY)	---
41		PRO	PRO	PRO	---	---	1	1	1(GLN)	---
42		LEU	LEU	LEU	---	---	1	1	1(ALA)	---
43		VAL	VAL	VAL	---	---	1	1	1(PRO)	---
44		ILE	ILE	ILE	---	---	1	1	1(LEU)	---
45		VAL	VAL	VAL	---	---	1	1	1(VAL)	---
46		ILE	ILE	ILE	---	---	1	1	1(ILE)	---
47		TYR	TYR	TYR	---	---	1	1	1(TYR)	---
48		GLY	GLY	GLY	---	---	1	1	1(GLY)	---
49		ARG	ARG	ARG	---	---	1	1	1(ARG)	---
50		ASN	ASN	ASN	---	---	1	1	1(ASN)	---
51		ASN	ASN	ASN	---	---	1	1	1(ASN)	---
52		ARG	ARG	ARG	---	---	1	1	1(ARG)	---
53		PRO	PRO	PRO	---	---	1	1	1(PRO)	---
54		SER	SER	SER	---	---	1	1	1(SER)	---
55		ILE	ILE	ILE	---	---	1	1	1(ILE)	---
56		PRO	PRO	PRO	---	---	1	1	1(PRO)	---
57		ASP	ASP	ASP	---	---	1	1	1(ASP)	---
58		ARG	ARG	ARG	---	---	1	1	1(ASP)	---
59		PHE	PHE	PHE	---	---	1	1	1(PHE)	---
60		SER	SER	SER	---	---	1	1	1(SER)	---
61		GLY	GLY	GLY	---	---	1	1	1(GLY)	---
62		SER	SER	SER	---	---	1	1	1(SER)	---
63		SER	SER	SER	---	---	1	1	1(SER)	---
64		SER	SER	SER	---	---	1	1	1(SER)	---
65		SER	SER	SER	---	---	1	1	1(SER)	---
66		SER	SER	SER	---	---	1	1	1(SER)	---
67		GLY	GLY	GLY	---	---	1	1	1(GLY)	---
68		HIS	HIS	HIS	---	---	1	1	1(HIS)	---
69		THR	THR	THR	---	---	1	1	1(THR)	---
70		ALA	ALA	ALA	---	---	1	1	1(ALA)	---
71		SER	SER	SER	---	---	1	1	1(SER)	---
72		LEU	LEU	LEU	---	---	1	1	1(LEU)	---
73		THR	THR	THR	---	---	1	1	1(THR)	---
74		ILE	ILE	ILE	---	---	1	1	1(ILE)	---
75		THR	THR	THR	---	---	1	1	1(THR)	---
76		GLY	GLY	GLY	---	---	1	1	1(GLY)	---
77		ALA	ALA	ALA	---	---	1	1	1(ALA)	---
78		GLN	GLN	GLN	---	---	1	1	1(GLN)	---
79		ALA	ALA	ALA	---	---	1	1	1(ALA)	---
80		GLU	GLU	GLU	---	---	1	1	1(GLU)	---
81		ASP	ASP	ASP	---	---	1	1	1(ASP)	---
82		GLU	GLU	GLU	---	---	1	1	1(GLU)	---
83		ALA	ALA	ALA	---	---	1	1	1(ALA)	---
84		ASP	ASP	ASP	---	---	1	1	1(ASP)	---
85		TYR	TYR	TYR	---	---	1	1	1(TYR)	---
86		CYS	CYS	CYS	---	---	1	1	1(CYS)	---
87		ASN	ASN	ASN	---	---	1	1	1(ASN)	---
88		ARG	ARG	ARG	---	---	1	1	1(ARG)	---
89		ASP	ASP	ASP	---	---	1	1	1(ASP)	---
90		SER	SER	SER	---	---	1	1	1(SER)	---
91		SER	SER	SER	---	---	1	1	1(SER)	---
92		SER	SER	SER	---	---	1	1	1(SER)	---
93		SER	SER	SER	---	---	1	1	1(SER)	---
94		GLY	GLY	GLY	---	---	1	1	1(GLY)	---
95		LYS	LYS	LYS	---	---	1	1	1(LYS)	---
95A		HIS	HIS	HIS	---	---	1	1	1(HIS)	---
95B		---	---	---	---	---	---	---	---	---
95C		---	---	---	---	---	---	---	---	---
95D		---	---	---	---	---	---	---	---	---
95E		---	---	---	---	---	---	---	---	---
95F		VAL	VAL	VAL	---	---	1	1	1(VAL)	---
96		LEU	LEU	LEU	---	---	1	1	1(LEU)	---
97		PHE	PHE	PHE	---	---	1	1	1(PHE)	---
98		GLY	GLY	GLY	---	---	1	1	1(GLY)	---
99		GLY	GLY	GLY	---	---	1	1	1(GLY)	---
100		GLY	GLY	GLY	---	---	1	1	1(GLY)	---
101		THR	THR	THR	---	---	1	1	1(THR)	---
102		LYS	LYS	LYS	---	---	1	1	1(LYS)	---
103		LEU	LEU	LEU	---	---	1	1	1(LEU)	---
104		THR	THR	THR	---	---	1	1	1(THR)	---
105		VAL	VAL	VAL	---	---	1	1	1(VAL)	---
106		LEU	LEU	LEU	---	---	1	1	1(LEU)	---
106A		GLY	GLY	GLY	---	---	1	1	1(GLY)	---
107		GLN	GLN	GLN	---	---	1	1	1(GLN)	---
108		PRO	PRO	PRO	---	---	1	1	1(PRO)	---

REFERENCE: HUMAN LAMBDA LIGHT CHAINS SUBGROUP IV

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NOTES: HUMAN LAMBDA LIGHT CHAINS SUBGROUP IV

IDENTICAL SETS OF FRAMEWORK SEGMENTS:

FR1: SET 1: SH11,NEV12; (2 IDENTICAL)

FR2:

FR3:

FR4: SET 1: SH11; (IDENTICAL TO 4 HUMAN V-LAMBDA-I: BL2 'CL16),RHE10),OKA12),NIG-51119); 5 HUMAN V-LAMBDA-II: MES12),ES492(8), TRG114),VIL117),WIN121); 4 HUMAN V-LAMBDA-III: NIL11),CAPI4),BAU112),DEL114); 3 HUMAN V-LAMBDA-VI: SU12),TNC14), LBV'CL181; AND 24 MOUSE V-LAMBDA: MQPC104E11),J558(2),X5104(3),HOPC114),J69(5),H2061(6),W3159(7),Y5431(8),Y5485(9), Y5930(10),Y566(9),11),MQPC511(L),12),S17(8),13),Y5444(14),Y5606(16),S17(8),16),H202(17),RPC2(8),18),IG 303(LAMBDA CL)19), S43 CL121),S2H5 CL13(8),S2E9 CL13(9),S1F12 CL14(40),IG 25(LAMBDA CL)411);

- THE FOLLOWING WERE EQUALLY AND MOST FREQUENTLY OCCURRING:

AT POSITION	RESIDUES
19	(VAL,ALA)
27	(LYS,SER)
30	(ALA,GLY,ASP,GLN)
32	(TYR,ASP,ASN)
34	(ILE,ALA,SER)

HUMAN LAMBDA LIGHT CHAINS SUBGROUP V

	INVARIENT RESIDUES	1 BO	2 HIS	3 MCGL	# OF SEQUENCES	# OF AMINO ACIDS	OCCURRENCES OF MOST COMMON AMINO ACID	VARIABILITY
F R 1	0	PCA	PCA	PCA	3	1	3(PCA)	1.
	1	SER	SER	SER	3	1	3(SER)	1.
	2	ALA	ALA	ALA	3	1	3(ALA)	1.
	3	LEU	LEU	LEU	3	1	3(LEU)	1.
	4	THR	THR	THR	3	1	3(THR)	1.
	5	GLN	GLN	GLN	3	1	3(GLN)	1.
	6	PRO	PRO	PRO	3	1	3(PRO)	1.
	7	PRO	PRO	PRO	3	1	3(PRO)	1.
	8	SER	SER	SER	3	1	3(SER)	1.
	9	---	---	---	3	1	---	1.
	10	ALA	ALA	ALA	3	1	3(ALA)	1.
	11	SER	SER	SER	3	1	3(SER)	1.
	12	GLY	GLY	GLY	3	1	3(GLY)	1.
	13	SER	SER	SER	3	1	3(SER)	1.
F R 2	14	---	---	---	3	1	---	1.
	15	PRO	PRO	PRO	3	2	2(PRO)	1.
	16	GLY	GLY	GLY	3	1	3(GLY)	1.
	17	GLN	GLN	GLN	3	1	3(GLN)	1.
	18	SER	SER	SER	3	1	3(SER)	1.
	19	VAL	VAL	VAL	3	1	3(VAL)	1.
	20	THR	THR	THR	3	1	3(THR)	1.
	21	ILE	ILE	ILE	3	1	3(ILE)	1.
	22	SER	SER	SER	3	1	3(SER)	1.
	23	CYS	CYS	CYS	3	1	3(CYS)	1.
	24	THR	THR	THR	3	1	3(THR)	1.
	25	GLY	GLY	GLY	3	1	3(GLY)	1.
	26	THR	THR	THR	3	1	3(THR)	1.
	27	SER	SER	SER	3	1	3(SER)	1.
F R 3	27A	---	---	---	3	1	---	1.
	27C	SER	SER	SER	3	1	3(SER)	1.
	27D	ASP	ASP	ASP	3	1	3(ASP)	1.
	27E	VAL	VAL	VAL	3	1	3(VAL)	1.
	27F	GLY	GLY	GLY	3	1	3(GLY)	1.
	28	---	---	---	3	1	---	1.
	29	ASP	ASP	ASP	3	1	3(ASP)	1.
	30	ASN	ASN	ASN	3	1	3(ASN)	1.
	31	LYS	LYS	LYS	3	1	3(LYS)	1.
	32	VAL	VAL	VAL	3	1	3(VAL)	1.
	33	THR	THR	THR	3	1	3(THR)	1.
	34	SER	SER	SER	3	1	3(SER)	1.
	35	TRP	TRP	TRP	3	1	3(TRP)	1.
	36	TYR	TYR	TYR	3	1	3(TYR)	1.
F R 4	37	GLN	GLN	GLN	3	1	3(GLN)	1.
	38	GLN	GLN	GLN	3	1	3(GLN)	1.
	39	HIS	HIS	HIS	3	1	3(HIS)	1.
	40	PRO	PRO	PRO	3	1	3(PRO)	1.
	41	GLY	GLY	GLY	3	1	3(GLY)	1.
	42	ARG	ARG	ARG	3	1	3(ARG)	1.
	43	ALA	ALA	ALA	3	1	3(ALA)	1.
	44	PRO	PRO	PRO	3	1	3(PRO)	1.
	45	LYS	LYS	LYS	3	1	3(LYS)	1.
	46	LEU	LEU	LEU	3	1	3(LEU)	1.
	47	VAL	VAL	VAL	3	1	3(VAL)	1.
	48	ILE	ILE	ILE	3	1	3(ILE)	1.
	49	PHE	PHE	PHE	3	1	3(PHE)	1.
	50	GLU	GLU	GLU	3	1	3(GLU)	1.
F R 5	51	VAL	VAL	VAL	3	1	3(VAL)	1.
	52	SER	SER	SER	3	1	3(SER)	1.
	53	GLY	GLY	GLY	3	1	3(GLY)	1.
	54	ARG	ARG	ARG	3	1	3(ARG)	1.
	55	PRO	PRO	PRO	3	1	3(PRO)	1.
	56	SER	SER	SER	3	1	3(SER)	1.
	57	GLY	GLY	GLY	3	1	3(GLY)	1.
	58	VAL	VAL	VAL	3	1	3(VAL)	1.
	59	PRO	PRO	PRO	3	1	3(PRO)	1.
	60	ASP	ASP	ASP	3	1	3(ASP)	1.
	61	ARG	ARG	ARG	3	1	3(ARG)	1.
	62	PHE	PHE	PHE	3	1	3(PHE)	1.
	63	SER	SER	SER	3	1	3(SER)	1.
	64	GLY	GLY	GLY	3	1	3(GLY)	1.
F R 6	65	SER	SER	SER	3	1	3(SER)	1.
	66	LYS	LYS	LYS	3	1	3(LYS)	1.
	67	SER	SER	SER	3	1	3(SER)	1.
	68	VAL	VAL	VAL	3	1	3(VAL)	1.
	69	ASN	ASN	ASN	3	1	3(ASN)	1.
	70	THR	THR	THR	3	1	3(THR)	1.
	71	ALA	ALA	ALA	3	1	3(ALA)	1.
	72	SER	SER	SER	3	1	3(SER)	1.
	73	LEU	LEU	LEU	3	1	3(LEU)	1.
	74	THR	THR	THR	3	1	3(THR)	1.
	75	VAL	VAL	VAL	3	1	3(VAL)	1.
	76	SER	SER	SER	3	1	3(SER)	1.
	77	GLY	GLY	GLY	3	1	3(GLY)	1.
	78	LEU	LEU	LEU	3	1	3(LEU)	1.
F R 7	79	ARG	ARG	ARG	3	1	3(ARG)	1.
	80	ALA	ALA	ALA	3	1	3(ALA)	1.
	81	GLU	GLU	GLU	3	1	3(GLU)	1.
	82	ASP	ASP	ASP	3	1	3(ASP)	1.
	83	GLU	GLU	GLU	3	1	3(GLU)	1.
	84	ALA	ALA	ALA	3	1	3(ALA)	1.
	85	ASP	ASP	ASP	3	1	3(ASP)	1.
	86	TYR	TYR	TYR	3	1	3(TYR)	1.
	87	TYR	TYR	TYR	3	1	3(TYR)	1.
	88	CYS	CYS	CYS	3	1	3(CYS)	1.
	89	SER	SER	SER	3	1	3(SER)	1.
	90	SER	SER	SER	3	1	3(SER)	1.
	91	TYR	TYR	TYR	3	1	3(TYR)	1.
	92	VAL	VAL	VAL	3	1	3(VAL)	1.
F R 8	93	ASP	ASP	ASP	3	1	3(ASP)	1.
	94	ASN	ASN	ASN	3	1	3(ASN)	1.
	95	ASN	ASN	ASN	3	1	3(ASN)	1.
	95A	---	---	---	3	1	---	1.
	95B	---	---	---	3	1	---	1.
	95C	---	---	---	3	1	---	1.
	95D	---	---	---	3	1	---	1.
	95E	---	---	---	3	1	---	1.
	95F	---	---	---	3	1	---	1.
	96	PHE	PHE	PHE	3	1	3(PHE)	1.
	97	VAL	VAL	VAL	3	1	3(VAL)	1.
	98	PHE	PHE	PHE	3	1	3(PHE)	1.
	99	GLY	GLY	GLY	3	1	3(GLY)	1.
	100	GLY	GLY	GLY	3	1	3(GLY)	1.
F R 9	101	GLY	GLY	GLY	3	1	3(GLY)	1.
	102	THR	THR	THR	3	1	3(THR)	1.
	103	LYS	LYS	LYS	3	1	3(LYS)	1.
	104	THR	THR	THR	3	1	3(THR)	1.
	105	VAL	VAL	VAL	3	1	3(VAL)	1.
	106	LEU	LEU	LEU	3	1	3(LEU)	1.
	107	ARG	ARG	ARG	3	1	3(ARG)	1.
	108	GLN	GLN	GLN	3	1	3(GLN)	1.
	109	PRO	PRO	PRO	3	1	3(PRO)	1.
	110	---	---	---	3	1	---	1.
	111	---	---	---	3	1	---	1.
	112	---	---	---	3	1	---	1.
	113	---	---	---	3	1	---	1.
	114	---	---	---	3	1	---	1.

ANTIBODY SPECIFICITIES: HUMAN LAMBDA LIGHT CHAINS SUBGROUP V

- 3) **MCG**: ANTI-EPSILON-DNP-LYS, EPSILON-DNP-AMINOCAPROATE, DNP-LEU, TRIACETIN, SODIUM MERTHIOLATE, METHADONE, 1,10-PHENANTHROLINE, CAFFEINE, THEOPHYLLINE, DI-DNP-LYS, DNP-TRP, DNP-PHE, DI-DNP-TYR, COLCHICINE, P-NITROANILINE, P-NITROPHENYLPHOSPHORYL CHOLINE, 5-ACETYLURACIL, MENADIONE, MEPERIDINE, TRIBUTYRIN, OMEGA-BROMOHEPTANOATE, O-CHLOROMERCURIPHENOL, P-CHLOROMERCURIPHENOL, PHENYLMERCURIC COMPOUNDS, METHYL-MERCURIC CHLORIDE.

REFERENCE: HUMAN LAMBDA LIGHT CHAINS SUBGROUP V

- 1) **BO**: WIKLER, M. & PUTNAM, F. W. (1970) J. BIOL. CHEM. 245, 4468-4507. (CHECKED BY AUTHOR 06/15/83)
 2) **HBJ2**: HOOD, L., GRAY, W. R., SANDERS, B. G. & DREYER, W. J. (1967) COLD SPRING HARBOR SYMP. QUANTITATIVE BIOL., 32, 133-145.
 3) **MCG**: FETT, J. W. & DEUTSCH, H. F. (1974) BIOCHEMISTRY 13, 4102-4114. (CHECKED BY AUTHOR)

NOTES: HUMAN LAMBDA LIGHT CHAINS SUBGROUP V**IDENTICAL SETS OF FRAMEWORK SEGMENTS:**

- FR1: SET 1: BO[1], HBJ2[2]; (2 IDENTICAL)
 FR2:
 FR3:
 FR4: SET 1: BO[1]; (IDENTICAL TO 1 HUMAN V-LAMBDA-I: NEWM[1]); AND 1 HUMAN V-LAMBDA-II: WH[3])
 SET 2: MCG[3]; (IDENTICAL TO 1 HUMAN V-LAMBDA-I: LOC[11])

IDENTICAL SETS OF COMPLEMENTARITY DETERMINING REGIONS

- CDR1: SET 1: MCG[3]; (IDENTICAL TO 2 HUMAN V-LAMBDA-II: MESI[2], VIL1[17])
 CDR2:
 CDR3:

- THE FOLLOWING WERE EQUALLY AND MOST FREQUENTLY OCCURRING:

AT POSITION	RESIDUES
29	(GLY, ASP)
30	(TYR, ASN)
31	(LYS, ASN)
40	(PRO, ALA)
42	(LYS, ARG)
46	(LEU, VAL)
47	(ILE, VAL)
49	(TYR, PHE)
52	(SER, ASN)
53	(LYS, GLY)
68	(GLY, ASP)
79	(ARG, GLN)
92	(VAL, GLU)
93	(GLY, ASP)
94	(SER, ASN)
95	(ASP, ASN)
100	(THR, GLY)
104	(LEU, VAL)
107	(ARG, GLY)

HUMAN LAMBDA LIGHT CHAINS SUBGROUP VI

[illegible]

[illegible]

HUMAN HEAVY CHAINS SUBGROUP 1 (cont'd)

	25*	26	27	28	29	30	31	32	33	34	# OF	# OF	OCCURRENCES	VARIABILITY
	KOH	RIC	WIS	LEU	SAC	DEE	LEA	HAR	HUS	SEQUENCES	AMINO ACIDS	OF MOST COMMON AMINO ACID		
F R 1	0	---	---	---	---	---	---	---	---	---	30	5	21(PCA)	7.1
	1	gln	PCA	PCA	PCA	gln	---	---	---	---	30	6	25(VAL)	7.2
	2	VAL	VAL	met	VAL	VAL	---	---	---	---	29	6	22(GLN)	7.3
	3	GLN	GLN	GLN	---	---	pea	pea	---	---	29	6	24(LEU)	7.1
	4	LEU	---	---	---	---	LEU	LEU	---	---	14	4	11(VAL)	5.1
	5	---	---	---	---	---	---	---	---	---	14	2	10(GLN)	2.8
	6	---	---	---	---	---	---	---	---	---	14	1	14(SER)	1
	7	---	---	---	---	---	---	---	---	---	14	1	14(GLY)	1
	8	---	---	---	---	---	---	---	---	---	15	4	12(ALA)	5
	9	---	---	---	---	---	---	---	---	---	14	3	12(GLU)	3.5
	10	---	---	---	---	---	---	---	---	---	14	3	12(VAL)	2.3
	11	---	---	---	---	---	---	---	---	---	15	5	9(LYS)	6.3
	12	---	---	---	---	---	---	---	---	---	14	2	13(LYS)	2.2
	13	---	---	---	---	---	---	---	---	---	14	2	13(PRO)	2.2
	14	---	---	---	---	---	---	---	---	---	14	3	12(GLY)	3.5
	15	---	---	---	---	---	---	---	---	---	12	4	4(+)	12
	16	---	---	---	---	---	---	---	---	---	11	2	10(SER)	2.2
	17	---	---	---	---	---	---	---	---	---	12	5	7(VAL)	8.6
	18	---	---	---	---	---	---	---	---	---	12	5	6(+)	6.5
	19	---	---	---	---	---	---	---	---	---	13	3	6(VAL)	8
C D R 1	20	---	---	---	---	---	---	---	---	---	12	4	6(VAL)	8
	21	---	---	---	---	---	---	---	---	---	11	3	8(SER)	3.7
	22	---	---	---	---	---	---	---	---	---	9	2	8(CYS)	2.3
	23	---	---	---	---	---	---	---	---	---	11	3	9(LYS)	3.7
	24	---	---	---	---	---	---	---	---	---	11	5	4(ALA)	14
	25	---	---	---	---	---	---	---	---	---	10	3	8(SER)	2.2
	26	---	---	---	---	---	---	---	---	---	10	4	5(TYR)	6
	27	---	---	---	---	---	---	---	---	---	10	3	6(THR)	4
	28	---	---	---	---	---	---	---	---	---	8	2	7(PHE)	2.3
	29	---	---	---	---	---	---	---	---	---	8	5	3(SER)	1.3
	30	---	---	---	---	---	---	---	---	---	8	7	2(ASP)	28
	31	---	---	---	---	---	---	---	---	---	8	6	5(TYR)	3.2
	32	---	---	---	---	---	---	---	---	---	8	6	2(+)	24
	33	---	---	---	---	---	---	---	---	---	8	4	4(ILE)	6
	34	---	---	---	---	---	---	---	---	---	8	5	3(HIS)	1.3
	35	---	---	---	---	---	---	---	---	---	8	2	7(TRP)	2.3
	36	---	---	---	---	---	---	---	---	---	8	3	5(VAL)	4.8
	37	---	---	---	---	---	---	---	---	---	8	2	7(ARG)	2.3
	38	---	---	---	---	---	---	---	---	---	8	2	7(GLN)	2.3
	39	---	---	---	---	---	---	---	---	---	8	3	6(ALA)	4
40	---	---	---	---	---	---	---	---	---	8	2	7(PRO)	2.3	
F R 2	41	---	---	---	---	---	---	---	---	---	8	2	7(GLY)	2.3
	42	---	---	---	---	---	---	---	---	---	7	4	2(+)	14
	43	---	---	---	---	---	---	---	---	---	7	1	7(GLY)	1
	44	---	---	---	---	---	---	---	---	---	7	1	7(LEU)	1
	45	---	---	---	---	---	---	---	---	---	7	1	7(GLU)	1
	46	---	---	---	---	---	---	---	---	---	7	1	7(TRP)	1
	47	---	---	---	---	---	---	---	---	---	7	2	4(VAL)	3.5
	48	---	---	---	---	---	---	---	---	---	7	2	6(GLY)	2.3
	49	---	---	---	---	---	---	---	---	---	7	3	1(+)	49
	50	---	---	---	---	---	---	---	---	---	7	6	2(ASN)	21
	51	---	---	---	---	---	---	---	---	---	7	6	5(ILE)	4.2
	52	---	---	---	---	---	---	---	---	---	6	3	4(PRO)	1
	53	---	---	---	---	---	---	---	---	---	7	6	2(SER)	21
	54	---	---	---	---	---	---	---	---	---	7	5	2(+)	18
	55	---	---	---	---	---	---	---	---	---	7	3	4(GLY)	5.3
	56	---	---	---	---	---	---	---	---	---	7	6	2(ASN)	21
	57	---	---	---	---	---	---	---	---	---	7	3	5(TYR)	4.2
	58	---	---	---	---	---	---	---	---	---	6	4	3(ALA)	6
	59	---	---	---	---	---	---	---	---	---	6	3	3(PRO)	6
	60	---	---	---	---	---	---	---	---	---	6	3	2(+)	12
C D R 2	61	---	---	---	---	---	---	---	---	---	6	3	3(PHE)	6
	62	---	---	---	---	---	---	---	---	---	6	3	4(GLN)	7
	63	---	---	---	---	---	---	---	---	---	7	1	3(GLY)	12
	64	---	---	---	---	---	---	---	---	---	7	4	4(THR)	1
	65	---	---	---	---	---	---	---	---	---	7	5	3(ASP)	2.7
	66	---	---	---	---	---	---	---	---	---	7	1	7(ARG)	1
	67	---	---	---	---	---	---	---	---	---	6	2	5(VAL)	2.4
	68	---	---	---	---	---	---	---	---	---	6	2	5(THR)	2.4
	69	---	---	---	---	---	---	---	---	---	7	3	3(+)	7
	70	---	---	---	---	---	---	---	---	---	7	3	4(THR)	3.5
	71	---	---	---	---	---	---	---	---	---	7	3	3(+)	7
	72	---	---	---	---	---	---	---	---	---	7	3	4(THR)	3.5
	73	---	---	---	---	---	---	---	---	---	7	1	7(SER)	1
	74	---	---	---	---	---	---	---	---	---	7	3	3(+)	7
	75	---	---	---	---	---	---	---	---	---	7	3	4(ASN)	5.3
	76	---	---	---	---	---	---	---	---	---	7	3	4(THR)	5.3
	77	---	---	---	---	---	---	---	---	---	7	3	4(ALA)	5.3
	78	---	---	---	---	---	---	---	---	---	7	3	5(TYR)	4.2
	79	---	---	---	---	---	---	---	---	---	7	2	6(MET)	2.3
	80	---	---	---	---	---	---	---	---	---	7	2	5(GLU)	4.2
F R 3	81	---	---	---	---	---	---	---	---	---	8	5	7(LEU)	2.3
	82	---	---	---	---	---	---	---	---	---	8	5	3(SER)	2.3
	82A	---	---	---	---	---	---	---	---	---	8	2	7(LEU)	4
	82B	---	---	---	---	---	---	---	---	---	8	4	4(ARG)	8
	82C	---	---	---	---	---	---	---	---	---	8	4	5(SER)	8
	83	---	---	---	---	---	---	---	---	---	8	4	5(GLU) : 4(GLU)	4.8
	84	---	---	---	---	---	---	---	---	---	8	1	8(ASP) : 7(ASP)	5
	85	---	---	---	---	---	---	---	---	---	8	2	6(THR)	4
	86	---	---	---	---	---	---	---	---	---	8	3	6(THR)	4
	87	---	---	---	---	---	---	---	---	---	8	3	6(VAL)	4
	88	---	---	---	---	---	---	---	---	---	8	3	6(TYR)	2.3
	89	---	---	---	---	---	---	---	---	---	8	3	6(TYR)	2.3
	90	---	---	---	---	---	---	---	---	---	9	2	8(CYS)	1
	91	---	---	---	---	---	---	---	---	---	9	3	6(ARG)	4.5
	92	---	---	---	---	---	---	---	---	---	9	2	2(+)	18
	93	---	---	---	---	---	---	---	---	---	9	5	2(TYR)	21
	94	---	---	---	---	---	---	---	---	---	7	6	2(GLY)	21
	95	---	---	---	---	---	---	---	---	---	7	5	2(PHE)	15
	96	---	---	---	---	---	---	---	---	---	6	5	3(SER)	15
	97	---	---	---	---	---	---	---	---	---	6	5	2(ASN)	15
98	---	---	---	---	---	---	---	---	---	5	4	2(ASP)	15	
99	---	---	---	---	---	---	---	---	---	5	4	2(ASP)	15	
100	---	---	---	---	---	---	---	---	---	4	4	2(TYR)	15	
C D R 3	100A	---	---	---	---	---	---	---	---	---	5	4	2(ASP)	15
	100B	---	---	---	---	---	---	---	---	---	5	4	2(ASP)	15
	100C	---	---	---	---	---	---	---	---	---	4	4	2(TYR)	15
	100D	---	---	---	---	---	---	---	---	---	4	4	1(+)	1
	100E	---	---	---	---	---	---	---	---	---	2	2	1(+)	1
	100F	---	---	---	---	---	---	---	---	---	2	2	1(+)	1
	100G	---	---	---	---	---	---	---	---	---	2	2	1(+)	1
	100H	---	---	---	---	---	---	---	---	---	1	1	1(TYR)	1
	100I	---	---	---	---	---	---	---	---	---	1	1	1(THR)	1
	100J	---	---	---	---	---	---	---	---	---	1	1	1(+)	1
	100K	---	---	---	---	---	---	---	---	---	3	3	1(+)	1
	101	---	---	---	---	---	---	---	---	---	7	4	3(ASP) : 2(+)	9.3
	102	---	---	---	---	---	---	---	---	---	6	5	3(TYR)	18
	103	---	---	---	---	---	---	---	---	---	8	2	6(THR)	2.7
	104	---	---	---	---	---	---	---	---	---	8	3	6(GLY)	4
	105	---	---	---	---	---	---	---	---	---	8	3	5(GLN) : 4(GLN)	4.8
	106	---	---	---	---	---	---	---	---	---	8	1	8(GLY)	1
	107	---	---	---	---	---	---	---	---	---	9	1	4(THR)	1
	108	---	---	---	---	---	---	---	---	---	8	3	6(LEU)	4
	109	---	---	---	---	---	---	---	---	---	8	3	6(VAL)	4
110	---	---	---	---	---	---	---	---	---	9	2	6(THR)	2.3	
111	---	---	---	---	---	---	---	---	---	11	1	9(VAL)	1	
112	---	---	---	---	---	---	---	---	---	10	2	9(SER)	2.2	
113	---	---	---	---	---	---	---	---	---	10	1	10(SER)	1	

ANTIBODY SPECIFICITIES: HUMAN HEAVY CHAINS SUBGROUP I

- 2) SIE: ANTI-HUMAN GAMMA G GLOBULIN; WA IDIOTYPE
- 4) WOL: ANTI-HUMAN GAMMA G GLOBULIN; WA IDIOTYPE
- 10) ST: COLD AGGLUTININ WITH ANTI-BLOOD GROUP I ACTIVITY
- 16) MAR: ANTI-LIPOPROTEIN LIPASE
- 25) KOH: ANTI-HUMAN GAMMA G GLOBULIN

CLASS: HUMAN HEAVY CHAINS SUBGROUP I

- 1) EU: IGG1-KAPPA
- 2) SIE: IGM-KAPPA
- 4) WOL: IGM-KAPPA
- 5) CA: IGG1-
- 6) ND'CL: IGE-
- 7) MOT: IGG-
- 8) BRO: IGG: IGG-KAPPA
- 10) STE: IGG1-
- 11) BEN(I): IGG3-
- 12) ZUC: IGG3-
- 13) DI: IGM-
- 14) BOT: IGM-
- 15) OMM'CL: IGG3-
- 16) MAR: IGM-
- 19) WAR: IGG1-
- 20) VL: IGG3-LAMBDA
- 21) UIN: IGG4-
- 22) ADA: IGA-
- 23) NOR: IGA-
- 24) SAW: IGM2-
- 25) KOH: IGM-LAMBDA
- 26) RIC: IGG3-
- 27) WIS: IGG3-
- 28) VAU: IGG1-
- 29) LEB: IGG1-
- 30) SAC: IGG1-KAPPA
- 34) HUS: IGG3-

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- 21) UIN: KAPLAN,A.P.,HOOD,L.,TERRY,W.D. & METZGER,H. (1971) IMMUNOCHEMISTRY,8,801-811. (CHECKED BY AUTHOR)
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NOTES: HUMAN HEAVY CHAINS SUBGROUP I

IDENTICAL SETS OF FRAMEWORK SEGMENTS:

- FR1: SET 1: VAU(26),LEB(29). (2 IDENTICAL)
- FR2: SET 1: EU(1),HGG(CU3). (2 IDENTICAL)
- SET 2: WOL(4). (IDENTICAL TO 2 HUMAN V-H-III: TIL(4),TE(10).)
- FR3: SET 1: ND(CU6). (IDENTICAL TO 1 HUMAN V-H-III: U266(CU106).)
- FR4: SET 1: WOL(4). (IDENTICAL TO 2 HUMAN V-H-III: MCE(4),NZU(15); 4 HUMAN V-H-III: TIL(4),DOB(3),WEA(33),NIE(34); AND 1 MOUSE V-H-III: MORC(2448).)
- SET 2: ND(CU6). (IDENTICAL TO 1 HUMAN V-H-III: HIG(CU10); 1 HUMAN V-H-III: U266(CU106); AND 1 MOUSE V-H-III: HOEX(1215).)

IDENTICAL SETS OF COMPLEMENTARY DETERMINING REGIONS:

- CDR1:
- CDR2:
- CDR3: SET 1: HGG(CU3). (IDENTICAL TO 1 HUMAN V-H-III: LAMBDA-VH26(CU2); 1 MOUSE V-H-III: PU14(CU22); AND 5 MOUSE V-H-III: 186-2(CU3), 186-1(CU5), 102(CU15), 23(CU18), 3(CU26).)
- SET 2: ND(CU6). (IDENTICAL TO 1 HUMAN V-H-III: U266(CU106).)

IDENTICAL SETS OF J-MINGENES:

- SET 1: ND(CU6). (IDENTICAL TO 1 HUMAN V-H-III: HIG(CU10); AND 1 HUMAN V-H-III: U266(CU106).)

NOTES: HUMAN HEAVY CHAINS SUBGROUP I (cont'd)

SPECIFIC NOTES:

- 3) **HQ2'CL**: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN FETAL LIVER GENOMIC DNA.
- 6) **ND'CL**: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA. IT CORRESPONDS TO THE AMINO ACID SEQUENCE DETERMINED EARLIER EXCEPT THAT THE AMINO ACID SEQUENCE DETERMINATION GAVE PCA AT POSITION 1, VAL AT 2, VAL AT 34, GLY AT 35, ILE AT 48 AND HIS AT 49.
- 7) **MOT**: PAPAINE CLEAVES BETWEEN ARG 56 AND THR 57, AND BETWEEN ARG 62 AND SER 63.
- 12) **ZUC**: IT WAS FROM A CASE OF HEAVY CHAIN DISEASE.
- 14) **BOT**: IT WAS FROM A CASE OF IGM HEAVY CHAIN DISEASE.
- 15) **OMM'CL**: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN CELL LINE CDNA. IT WAS FROM A CASE OF HEAVY CHAIN DISEASE.
- 27) **WIS**: IT WAS FROM A CASE OF HEAVY CHAIN DISEASE. ITS RESIDUES AT POSITIONS 108 AND 109 ARE ASN AND CYS RESPECTIVELY, WHICH DO NOT CORRESPOND TO THE USUAL RESIDUES FOUND AT THESE POSITIONS IN HUMAN HEAVY CHAIN SUBGROUP I.
- 28) **VAU**: IT WAS FROM A CASE OF HEAVY CHAIN DISEASE.
- 29) **LEB**: IT WAS FROM A CASE OF HEAVY CHAIN DISEASE.
- 30) **SAC**: IT WAS FROM A CASE OF HEAVY CHAIN DISEASE.

~ THE FOLLOWING WERE EQUALLY AND MOST FREQUENTLY OCCURRING:

AT. POSITION	RESIDUES
16	(ALA,SER)
19	(LYS,ARG)
33	(TYR,ALA)
43	(LYS,ARG,GLN)
50	(TRP,ILE,VAL,SER,GLY,GLU,GLN)
54	(PHE,SER)
56	(PRO,GLY)
62	(LYS,ARG)
69	(VAL,MET)
71	(LEU,ARG)
73	(PRO,THR)
75	(PHE,THR)
85	(GLY,GLU)
100D	(TYR,PRO,SER,ASN)
100E	(PHE,GLY)
100F	(THR,ASP)
100G	(TYR,SER)
100H	(LEU,SER)
100K	(TYR,PHE,LEU)
101	(PRO,ASP)

HUMAN HEAVY CHAINS SUBGROUP I

	INVIANT RESIDUES	COR	2	OU	MCE	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	100A	100B	100C	100D	100E	100F	100G	100H	100I	100J	100K	100L	100M	100N	100O	100P	100Q	100R	100S	100T	100U	100V	100W	100X	100Y	100Z	100AA	100AB	100AC	100AD	100AE	100AF	100AG	100AH	100AI	100AJ	100AK	100AL	100AM	100AN	100AO	100AP	100AQ	100AR	100AS	100AT	100AU	100AV	100AW	100AX	100AY	100AZ	100BA	100BB	100BC	100BD	100BE	100BF	100BG	100BH	100BI	100BJ	100BK	100BL	100BM	100BN	100BO	100BP	100BQ	100BR	100BS	100BT	100BU	100BV	100BW	100BX	100BY	100BZ	100CA	100CB	100CC	100CD	100CE	100CF	100CG	100CH	100CI	100CJ	100CK	100CL	100CM	100CN	100CO	100CP	100CQ	100CR	100CS	100CT	100CU	100CV	100CW	100CX	100CY	100CZ	100DA	100DB	100DC	100DD	100DE	100DF	100DG	100DH	100DI	100DJ	100DK	100DL	100DM	100DN	100DO	100DP	100DQ	100DR	100DS	100DT	100DU	100DV	100DW	100DX	100DY	100DZ	100EA	100EB	100EC	100ED	100EE	100EF	100EG	100EH	100EI	100EJ	100EK	100EL	100EM	100EN	100EO	100EP	100EQ	100ER	100ES	100ET	100EU	100EV	100EW	100EX	100EY	100EZ	100FA	100FB	100FC	100FD	100FE	100FF	100FG	100FH	100FI	100FJ	100FK	100FL	100FM	100FN	100FO	100FP	100FQ	100FR	100FS	100FT	100FU	100FV	100FW	100FX	100FY	100FZ	100GA	100GB	100GC	100GD	100GE	100GF	100GG	100GH	100GI	100GJ	100GK	100GL	100GM	100GN	100GO	100GP	100GQ	100GR	100GS	100GT	100GU	100GV	100GW	100GX	100GY	100GZ	100HA	100HB	100HC	100HD	100HE	100HF	100HG	100HH	100HI	100HJ	100HK	100HL	100HM	100HN	100HO	100HP	100HQ	100HR	100HS	100HT	100HU	100HV	100HW	100HX	100HY	100HZ	100IA	100IB	100IC	100ID	100IE	100IF	100IG	100IH	100II	100IJ	100IK	100IL	100IM	100IN	100IO	100IP	100IQ	100IR	100IS	100IT	100IU	100IV	100IW	100IX	100IY	100IZ	100JA	100JB	100JC	100JD	100JE	100JF	100JG	100JH	100JI	100JJ	100JK	100JL	100JM	100JN	100JO	100JP	100JQ	100JR	100JS	100JT	100JU	100JV	100JW	100JX	100JY	100JZ	100KA	100KB	100KC	100KD	100KE	100KF	100KG	100KH	100KI	100KJ	100KK	100KL	100KM	100KN	100KO	100KP	100KQ	100KR	100KS	100KT	100KU	100KV	100KW	100KX	100KY	100KZ	100LA	100LB	100LC	100LD	100LE	100LF	100LG	100LH	100LI	100LJ	100LK	100LM	100LN	100LO	100LP	100LQ	100LR	100LS	100LT	100LU	100LV	100LW	100LX	100LY	100LZ	100MA	100MB	100MC	100MD	100ME	100MF	100MG	100MH	100MI	100MJ	100MK	100ML	100MN	100MO	100MP	100MQ	100MR	100MS	100MT	100MU	100MV	100MW	100MX	100MY	100MZ	100NA	100NB	100NC	100ND	100NE	100NF	100NG	100NH	100NI	100NJ	100NK	100NL	100NM	100NO	100NP	100NQ	100NR	100NS	100NT	100NU	100NV	100NW	100NX	100NY	100NZ	100OA	100OB	100OC	100OD	100OE	100OF	100OG	100OH	100OI	100OJ	100OK	100OL	100OM	100ON	100OO	100OP	100OQ	100OR	100OS	100OT	100OU	100OV	100OW	100OX	100OY	100OZ	100PA	100PB	100PC	100PD	100PE	100PF	100PG	100PH	100PI	100PJ	100PK	100PL	100PM	100PN	100PO	100PP	100PQ	100PR	100PS	100PT	100PU	100PV	100PW	100PX	100PY	100PZ	100QA	100QB	100QC	100QD	100QE	100QF	100QG	100QH	100QI	100QJ	100QK	100QL	100QM	100QN	100QO	100QP	100QQ	100QR	100QS	100QT	100QU	100QV	100QW	100QX	100QY	100QZ	100RA	100RB	100RC	100RD	100RE	100RF	100RG	100RH	100RI	100RJ	100RK	100RL	100RM	100RN	100RO	100RP	100RQ	100RR	100RS	100RT	100RU	100RV	100RW	100RX	100RY	100RZ	100SA	100SB	100SC	100SD	100SE	100SF	100SG	100SH	100SI	100SJ	100SK	100SL	100SM	100SN	100SO	100SP	100SQ	100SR	100SS	100ST	100SU	100SV	100SW	100SX	100SY	100SZ	100TA	100TB	100TC	100TD	100TE	100TF	100TG	100TH	100TI	100TJ	100TK	100TL	100TM	100TN	100TO	100TP	100TQ	100TR	100TS	100TT	100TU	100TV	100TW	100TX	100TY	100TZ	100UA	100UB	100UC	100UD	100UE	100UF	100UG	100UH	100UI	100UJ	100UK	100UL	100UM	100UN	100UO	100UP	100UQ	100UR	100US	100UT	100UU	100UV	100UW	100UX	100UY	100UZ	100VA	100VB	100VC	100VD	100VE	100VF	100VG	100VH	100VI	100VJ	100VK	100VL	100VM	100VN	100VO	100VP	100VQ	100VR	100VS	100VT	100VU	100VV	100VW	100VX	100VY	100VZ	100WA	100WB	100WC	100WD	100WE	100WF	100WG	100WH	100WI	100WJ	100WK	100WL	100WM	100WN	100WO	100WP	100WQ	100WR	100WS	100WT	100WU	100WV	100WW	100WX	100WY	100WZ	100XA	100XB	100XC	100XD	100XE	100XF	100XG	100XH	100XI	100XJ	100XK	100XL	100XM	100XN	100XO	100XP	100XQ	100XR	100XS	100XT	100XU	100XV	100XW	100XX	100XY	100XZ	100YA	100YB	100YC	100YD	100YE	100YF	100YG	100YH	100YI	100YJ	100YK	100YL	100YM	100YN	100YO	100YP	100YQ	100YR	100YS	100YT	100YU	100YV	100YW	100YX	100YY	100YZ	100ZA	100ZB	100ZC	100ZD	100ZE	100ZF	100ZG	100ZH	100ZI	100ZJ	100ZK	100ZL	100ZM	100ZN	100ZO	100ZP	100ZQ	100ZR	100ZS	100ZT	100ZU	100ZV	100ZW	100ZX	100ZY	100ZZ
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HUMAN HEAVY CHAINS SUBGROUP II (cont'd)

VARIABILITY

	0	
	1	4.5
	2	5.3
	3	8
	4	2.2
	5	11
	6	2.2
	7	12
	8	3.7
	9	2.2
	10	7.5
	11	1
	12	1
	13	2.5
	14	2.2
F	15	6
R	16	6
1	17	2.2
	18	1
	19	3.3
	20	1
	21	1
	22	1
	23	3.6
	24	6
	25	2.2
	26	1
	27	14
	28	3.7
	29	6
	30	5.7
C	31	13
D	32	36
R	33	6
1	34	10
	35	40
	35A	
	35B	
	36	1
	37	2.2
	38	1
	39	2.2
	40	3.6
F	41	1
R	42	1
2	43	5
	44	4
	45	1
	46	1
	47	1
	48	3.3
	49	3.3
	50	23
	51	6.7
	52	20
	52A	
	52B	
	52C	
	53	11
C	54	10
D	55	13
R	56	3.3
2	57	8
	58	15
	59	6.7
	60	5.7
	61	17
	62	20
	63	2.2
	64	1
	65	7.5
	66	6
	67	1
	68	3.3
	69	3.3
	70	6.7
	71	4.3
	72	8
	73	2.2
	74	2.2
	75	1
	76	2.9
	77	1
	78	2.3
	79	3.3
F	80	1
R	81	20
3	82	10
	82A	
	82B	
	82C	
	83	11
	84	3.1
	85	6.6
	86	1
	87	2.4
	88	2.4
	89	4.7
	90	1
	91	2.4
	92	1
	93	2.2
	94	2.8
	95	26
	96	50
	97	19
	98	26
	99	35
	100	40
	100A	
C	100B	
D	100C	
R	100D	
3	100E	
	100F	
	100G	
	100H	
	100I	
	100J	
	100K	
	101	3.6
	102	6.3
	103	2.2
	104	2.2
	105	6
F	106	2.2
R	107	2.2
4	108	3.3
	109	12
	110	5.3
	111	2.2
	112	2.2
	113	4

ANTIBODY SPECIFICITIES: HUMAN HEAVY CHAIN γ GROUP II

8) NEWM: ANTI-3-(3-HYDROXY-3,7,11,15-TETRAMETHYL HEXADECYL) 2-METHYL 1,4 NAPHTHOQUINONE(VIT.K10H)

CLASS: HUMAN HEAVY CHAINS SUBGROUP II

- 1) COR: IGG1-
 2) DAW: IGG1-LAMBDA
 3) QU: IGM-KAPPA
 4) MCE: IGM-KAPPA
 6) HE: IGG1-
 8) NEWM: IGG1-LAMBDA
 9) WAH: IGG-LAMBDA
 12) SA: IGG2-LAMBDA
 15) NZU: IGM-
 16) ERI: IGD-

REFERENCE: HUMAN HEAVY CHAINS SUBGROUP II

- 1) COR: PRESS.E.M. & HOGG.N.M. (1970) BIOCHEM.J.117,641-660. (CHECKED BY AUTHOR)
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 4) MCE: GERBER-JENSON.B.,KAZIW.A.,KEHOE.J.M.,SCHEFFEL.C.,ERICKSON.B.W. & LITMAN.G.W. (1981) J.IMMUNOL.126,1212-1216. (CHECKED BY AUTHOR 12/19/80)
 5) CE-1 CL: TAKAHASHI.N.,NOMAT.T. & HONJO.T. (1984) PROC.NAT.ACAD.SCI.USA81,5194-5198.
 6) HE: CUNNINGHAM.B.A.,GOTTLEB.P.D.,PFLUM.M.N. & EDELMAN.G.M. (1971) PROGRESS IN IMMUNOLOGY (B.AMOS.ED.),ACADEMIC PRESS.N.Y.,PP.3-24. (CHECKED BY AUTHOR)
 7) SUP-T1 VH-JA-CL: DENNY.C.T.,YOSHIKAWA.M.,MAK.T.W.,SMITH.S.D.,HOLLIS.G.F. & KIRSCH.I.R. (1986) NATURE,320,548-551.
 8) NEWM: POLJAK.R.J.,AMZELL.L.M.,CHEN.B.L.,PHIZACKERLEY.R.P. & SAUL.F. (1974) PROC.NAT.ACAD.SCI.USA71,3440-3444. (CHECKED BY AUTHOR WHO CORRECTED RESIDUES 6,8-16,18,24,26,27,29 THROUGH 36,59,60 AS GIVEN IN TABLE OF THE FIRST EDITION OF THIS BOOK, AND HAS MORE RECENTLY REVISED RESIDUES 5,24,28,29,30,31,33,34,35,36,35,36,59,60 AND 101); POLJAK.R.J.,AMZELL.L.M.,CHEN.B.L.,CHU.Y.Y.,PHIZACKERLEY.R.P. & SAUL.F. (1976) COLD SPRING HARBOR SYMP. QUANTITATIVE BIOL. 41,639-645; POLJAK.R.J.,NAKASHIMA.Y.,CHEN.B.L. & KONIGSBERG.W. (1977) BIOCHEMISTRY,16,3412-3420. THE SEQUENCE LISTED IN THE LAST REFERENCE IS GIVEN IN THE TABLE. (CHECKED BY AUTHOR 11/30/81)
 9) WAH: PUTNAM.F.W.,TAKAHASHI.N.,TETAERT.D.,DEBUIRE.B. & LIN.L. (1981) PROC.NAT.ACAD.SCI.USA78,6168-6172. (CHECKED BY AUTHOR 11/30/81); TAKAHASHI.N.,TETAERT.D.,DEBUIRE.B.,LIN.L. & PUTNAM.F.W. (1982) PROC.NAT.ACAD.SCI.USA79,2850-2854.
 10) HIG1 CL: KUJO.A.,ISHIMIZU.T.,MISHIMURA.Y. & WATANABE.T. (1985) GENE,33,181-189. (CHECKED BY AUTHOR 10/01/85)
 11) CAR: FRANGIONE.B. (1966) PH.D. THESIS, UNIVERSITY OF CAMBRIDGE. (CHECKED BY AUTHOR)
 12) SA: MILSTEIN.C. & FRANGIONE.B. (1971) BIOCHEM.J.121,217-225. (CHECKED BY AUTHOR)
 13) ID: MONTGOMERY.P.C.,BELLO.A.C. & ROCKEY.J.H. (1970) BIOCHIM.BIOPHYS.ACTA,200,258-266. (CHECKED BY AUTHOR)
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 15) NZU: ERICKSON.B.W.,GERBER-JENSON.B.,WANG.A.C. & LITMAN.G.W. (1981) MOL.IMMUNOL.19,357-365. (CHECKED BY AUTHOR 11/30/81)
 16) ERI: MILSTEIN.C.P. & DEVERSON.E.V. (1980) IMMUNOLOGY,40,557-664. (CHECKED BY AUTHOR 11/30/82)

NOTES: HUMAN HEAVY CHAINS SUBGROUP II

IDENTICAL SETS OF FRAMEWORK SEGMENTS:

FR1:

FR2: SET 1: SUP-T1 VH-JA-CL7;WAH(9). (2 IDENTICAL)

FR3:

FR4: SET 1: MCE(4);NZU(15). (2 IDENTICAL HUMAN V-H-II; ALSO 1 HUMAN V-H-I: WO(4); 4 HUMAN V-H-III: TI(4);DOBI(3);WEA(33);NIE(34);

SET 2: HIG1-CL10(1) (IDENTICAL TO 1 HUMAN V-H-I: ND(4); 1 HUMAN V-H-III: U266(4); 1 MOUSE V-H-III: HDX12(15).)

IDENTICAL SETS OF J-MINIGENES:

SET 1: HIG1-CL10(1) (IDENTICAL TO 1 HUMAN V-H-I: ND(4); AND 1 HUMAN V-H-III: U266(4); 106(1))

SPECIFIC NOTES:

4) MCE: IT IS A CRYOIMMUNOGLOBULIN AND IS DESIGNATED BY THE AUTHORS AS MCE. IN ORDER TO DIFFERENTIATE IT FROM ANOTHER MCE SEQUENCED BY CAPRA ET AL., IT IS DENOTED AS MCE.

5) CE-1 CL: CELL LINE CESS

7) SUP-T1 VH-JA-CL: IT IS FROM A PATIENT SUFFERING FROM CHILDHOOD T-CELL LYMPHOMA WITH INV(4)(q11.2;q32.2). THE INVERSION ON CHROMOSOME 14 BRINGS THE VH GENE AND JA MINIGENE TOGETHER, GIVING RISE TO A HYBRID MOLECULE CONTAINING PART OF THE IMMUNOGLOBULIN GENE AND PART OF THE T-LYMPHOCYTE RECEPTOR FOR ANTIGEN GENE.

14) SPA: IT WAS FROM A CASE OF HEAVY CHAIN DISEASE.

15) NZU: IT IS A CRYOIMMUNOGLOBULIN.

- THE FOLLOWING WERE EQUALLY AND MOST FREQUENTLY OCCURRING:

AT POSITION	RESIDUES
5	(ARG, GLN)
10	(ALA, GLY)
32	(THR, SER, ASP)
35	(CY, SER)
44	(ALA, GLY)
52A	(TYR, HIS)
59	(SER, ASN)
81	(LYS, THR)
82	(LEU, MET)
82A	(THR, SER)
82B	(SER, ASN)
82C	(VAL, MET)
85	(VAL, ALA)
96	(PRO, LEU)
99	(PRO, ARG, GLY)
100A	(TYR, PHE)
100B	(ALA, THR)
100D	(TYR, LEU)
100F	(TYR, GLY)
100H	(TYR, SER, ASP, ASN)
100I	(SER, GLY, ASP)

[illegible]